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EVALUATION OF TEARING ENERGIES IN
STYRENE-BUTADIENE RUBBERS

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**EVALUATION OF TEARING ENERGIES IN
STYRENE-BUTADIENE RUBBERS***

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ABSTRACT

Tearing energies were obtained on samples removed from commercial tank-track pads and on a series of SBR laboratory formulations in which the carbon-black content varied from 0 to 40 wt.%. The commercial materials were tested at a strain rate of 10^{-1} /s with temperatures ranging from 22 to 140° C. The laboratory materials were tested also at both lower and higher strain rates and at -20° C. We evaluated the effect of temperature, strain rate, and carbon-black content on tearing energy, failure stress, and failure strain. The results, including the shape of the stress-strain curves, are discussed in terms of viscoelasticity, uncoiling of the long molecular chains, carbon-black rubber interactions, and knotty tearing.

In the absence of knotty tearing and for carbon-black contents of 20 wt.% or less the tearing energy, the failure stress, and failure elongation increased with either a decrease in temperature or an increase in carbon-black content. With the first appearance of knotty tearing, which occurred at 20 to 25 wt.% carbon black, large increases in tearing energy and failure strength were obtained. Between 25 and 40 wt.% carbon black both tearing energy and failure stress varied irregularly with either the carbon-black content or temperature. The

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trends obtained with the commercial materials are consistent with those exhibited by the laboratory formulations in the 25 to 40% carbon-black range. The results for the 35-wt.% formulation fell within the range of results obtained for the commercial materials.

TABLE OF CONTENTS

	Page
ABSTRACT	1
LIST OF TABLES.....	iv
LIST OF FIGURES.....	v
1. INTRODUCTION.....	1
2. EXPERIMENTAL.....	2
2.1 Test Materials.....	2
2.2 Test Procedure.....	3
2.3 Evaluation of Tearing Energy.....	4
3. RESULTS.....	5
3.1 Tearing-Energy Data.....	5
3.2 Stress-Strain Curves at 0.33 in./s.....	6
3.3 Failure-Stress Plots.....	8
3.4 Deviated, Knotty Tearing.....	8
4. DISCUSSION.....	10
5. SUMMARY.....	16
6. RECOMMENDATIONS FOR FURTHER STUDY.....	18
7. ACKNOWLEDGMENTS.....	19
8. REFERENCES.....	19

LIST OF TABLES

	Page
Table 1. Designations of Elastomers Tested.....	21
Table 2. Tearing Energy Data for SBR-0.....	22
Table 3. Tearing Energy Data for SBR-15.....	23
Table 4. Tearing Energy Data for SBR-20. All Tests Performed at a Crosshead Rate of 0.33 in./s (8.33 mm/s).....	24
Table 5. Tearing Energy Data for SBR-25. All Tests Performed at a Crosshead Rate of 0.33 in./s (8.33 mm/s).....	25
Table 6. Tearing Energy Data for SBR-30 All Tests Performed at a Crosshead Rate of 0.33 in./s (8.33 mm/s).....	25
Table 7. Tearing Energy Data for SBR-35.....	26
Table 8. Tearing Energy Data for SBR-40. All Tests Performed at a Crosshead Rate of 0.33 in./s (8.33 mm/s).....	27
Table 9. Tearing Energy Data for T142 Goodyear Pads-115 and -145 and T156 Goodyear Pad. All Tests Performed at a Crosshead Rate of 0.33 in./s (8.33 mm/s).....	28
Table 10. Tearing Energy Data for T142 Firestone Pad-117 and T156 Firestone Pad. All Tests Performed at a Crosshead rate of 0.33 in./s (8.33 mm/s).....	29
Table 11. Tearing Energy Data for T156 Standard Pad and T156 Standard Pad-380. All Tests Performed at a Crosshead Rate of 0.33 in./s (8.33 mm/s).....	30
Table 12. Relative Ratings on the Extent of Deviated, Knotty Tearing.....	31

LIST OF FIGURES

- Fig. 1 - Tearing energy as a function of temperature for a series of laboratory-formulated SBR elastomers tested at a crosshead rate of 0.33 in./s ($\dot{\epsilon} \approx 10^{-1}/s$).
- Fig. 2 - Tearing energy as a function of temperature for samples removed from tank track pads. Samples tested at a crosshead rate of 0.33 in./s ($\dot{\epsilon} \approx 10^{-1}/s$). Results for SBR-35 are included.
- Fig. 3 - Tearing energy as a function of temperature for three laboratory-formulated SBR elastomers tested at a high stroke-displacement rate of approx. 235 in./s ($\dot{\epsilon} \approx 470/s$ for un-nicked samples).
- Fig. 4 - Tearing energy as a function of carbon-black content. Samples tested at a stroke rate of 0.33 in./s ($\dot{\epsilon} \approx 10^{-1}/s$) at five temperatures.
- Fig. 5 - Tearing energy as a function of carbon-black content. Samples tested at a stroke rate of 235 in./s ($\dot{\epsilon} \approx 470/s$ for un-nicked samples) at four temperatures.
- Fig. 6 - Effect of strain rate on tearing energy at various temperatures for SBR-0, -15, and -35. (Strain rates are for un-nicked samples).
- Fig. 7 - Stress-strain curves for un-nicked SBR-0 strip samples tested at various temperatures at a crosshead rate of 0.33 in./s ($\dot{\epsilon} \approx 10^{-1}/s$). Insert plotted on same scale used for stress-strain curves of remaining formulations.
- Fig. 8 - Stress-strain curves for un-nicked SBR-15 strip samples tested at various temperatures at a crosshead rate of 0.33 in./s ($\dot{\epsilon} \approx 10^{-1}/s$).
- Fig. 9 - Stress-strain curves for un-nicked SBR-20 strip samples tested at various temperatures at a crosshead rate of 0.33 in./s ($\dot{\epsilon} \approx 10^{-1}/s$).
- Fig. 10 - Stress-strain curves for un-nicked SBR-25 strip samples tested at various temperatures at a crosshead rate of 0.33 in./s ($\dot{\epsilon} \approx 10^{-1}/s$).
- Fig. 11 - Stress-strain curves for un-nicked SBR-30 strip samples tested at various temperatures at a crosshead rate of 0.33 in./s ($\dot{\epsilon} \approx 10^{-1}/s$).
- Fig. 12 - Stress-strain curves for un-nicked SBR-35 strip samples tested at various temperatures at a crosshead rate of 0.33 in./s ($\dot{\epsilon} \approx 10^{-1}/s$).

LIST OF FIGURES (continued)

- Fig. 13 - Stress-strain curves for un-nicked SBR-40 strip samples tested at various temperatures at a crosshead rate of 0.33 in./s ($\dot{\epsilon} = 10^{-1}/s$).
- Fig. 14 - Stress-strain curves for un-nicked SBR strip samples tested at -20° C at a crosshead rate of 0.33 in./s ($\dot{\epsilon} = 10^{-1}/s$).
- Fig. 15 - Stress strain curves for un-nicked SBR strip samples tested at 22° C at a crosshead rate of 0.33 in./s ($\dot{\epsilon} = 10^{-1}/s$).
- Fig. 16 - Stress-strain curves for un-nicked SBR strip samples tested at 60° C at a crosshead rate of 0.33 in./s ($\dot{\epsilon} = 10^{-1}/s$).
- Fig. 17 - Stress-strain curves for un-nicked SBR strip samples tested at 100° C at a crosshead rate of 0.33 in./s ($\dot{\epsilon} = 10^{-1}/s$).
- Fig. 18 - Stress-strain curves for un-nicked SBR strip samples tested at 140° C at a crosshead rate of 0.33 in./s ($\dot{\epsilon} = 10^{-1}/s$).
- Fig. 19 - Failure stress of nicked SBR strip samples tested at various temperatures at a crosshead rate of 0.33 in./s ($\dot{\epsilon} = 10^{-1}/s$).
- Fig. 20 - Failure stress of un-nicked SBR strip samples tested at various temperatures at a crosshead rate of 0.33 in./s ($\dot{\epsilon} = 10^{-1}/s$).
- Fig. 21 - Failure stress of nicked SBR strip samples as a function of carbon-black content at a series of temperatures. Tested at a crosshead rate of 0.33 in./s ($\dot{\epsilon} = 10^{-1}/s$).
- Fig. 22 - Failure stress of un-nicked SBR strip samples as a function of carbon-black content at a series of temperatures. Tested at a crosshead rate of 0.33 in./s ($\dot{\epsilon} = 10^{-1}/s$).
- Fig. 23 - Effect of strain rate on failure stress at various temperatures for nicked strip samples of SBR-0, -15, and -35.
- Fig. 24 - Effect of strain rate on failure stress or maximum recorded stress at various temperatures for un-nicked strip samples of SBR-0, -15, and -35. NOTE: A number of the SBR-15 and SBR-35 samples could not be taken to failure at the high strain rate.
- Fig. 25 - Effect of filler content on tensile strain at inflection point along the engineering stress-strain curves obtained at a crosshead rate of 0.33 in./s over the temperature range of -20 to 140° C. Curve A refers to experimental data with points representing average values and bars the spread in values; Curve B refers to the effect of rubber phase dilution; Curve C refers to filler-rubber interactions; Curve D refers to a minimum observed inflection strain for SBR-0; $Z = X + Y$.

1. INTRODUCTION

Relative to the reliability of other tank components, the performance of tank track pads over most types of terrains is far from acceptable. The frequent premature wear, fracturing, and chunking of these pads result in costs amounting to several hundred million dollars annually. As part of an overall program supported by the U.S. Army, we have been involved in evaluating mechanical properties and deformation behavior of a number of styrene-butadiene polymeric elastomers, which are loaded with various amounts of carbon-black filler. Both laboratory-prepared formulations and samples cut out of commercial tank pads were evaluated. In this paper we report primarily on the tearing energies and failure stresses obtained for these materials tested at various temperatures and strain rates. We also include stress-strain curves for a number of the laboratory formulations and some discussion on strength and tearing behavior.

As first formulated by Rivlin and Thomas⁽¹⁾ the tearing energy can be described in terms of the strain-energy release rate of an advancing tear, in that the driving force for the tearing is provided by the release of stored elastic strain energy. The tearing becomes critical, i.e., self-propagating, when the decrease in this energy is equal to or greater than the work required to propagate the tear.

Various sample configurations have been used to evaluate the tearing energy, with the results virtually being independent of the type of test and/or sample configuration used.^(2,3,4) The derivations all consider that the overall strain-energy release rate is closely related to the strain-energy density at the tear tip. In our studies we elected to evaluate tearing energies using double-edge-nicked flat samples.

In a previous report⁽⁵⁾ we discussed in some detail the tearing behavior of a number of SBR formulations tested under a wide range of temperatures and strain rates. Included in that report were evaluations of torn and fractured surfaces of many of the samples listed in the present paper. Of special interest were the initiation and propagation

of tears that approached directions parallel to the tensile axis. These tear deviations have been ascribed to a strength-anisotropy phenomenon⁽⁶⁾ that is developed in the deformation of carbon-black-loaded elastomers tested under certain conditions of temperature and strain rate. This deviated tearing is frequently referred to as "knotty tearing" because of the knot-like features seen on the torn surfaces. It has been suggested that the blunting of the tear front, resulting from this deviated tearing, causes an increase in tensile strength and tearing energy of the elastomer, adding to the strengthening effect imparted by the carbon black, per se.^(2,6,7,8) In the present paper we examine the role of this phenomenon in attempting to relate our calculated tearing energies to the stress-strain curves of the un-nicked corresponding samples to the tensile strength of both nicked and un-nicked samples, and to the effect of strain rate, temperature, and carbon-black loadings.

2. EXPERIMENTAL

2.1 Test Materials

The materials evaluated are listed in Table 1. Styrene-butadiene rubber (SBR) is the polymeric component in both types of materials. The carbon-black content of the laboratory formulations ranges from 0 to 40 wt. %; the 35-wt. % content is close to that contained in commercial tank track pads. Most of the laboratory formulations were compounded and cured by Burke Industries, San Jose, CA. The materials were obtained in the form of 6 in. x 6 in. (152 mm x 152 mm) sheets, ranging in thickness from 0.070 in. to 0.095 in. (1.78 to 2.41 mm). In the later stages of the program the test materials were made in our own laboratory, providing sheets measuring 10 in. (254 mm) diameter x 0.080 in. (2.03 mm) thick. Monsanto Rheometer tests for t_{90} (time to reach 90% of maximum torque) of our laboratory-compounded materials were made by Harold C. Seger, Rubber Consultant, Emeryville, CA.

Test samples of commercial materials (Goodyear, Firestone, and Standard T142 and T156 pads) were obtained from sheets that were sawed off from the actual pads. The sheets were chilled in a freon-dry ice mixture and machined to a smooth finish giving a thickness of approximately 0.100 in. (2.54 mm), the sheets being re-chilled between successive cuts. The individual sheets from a pad are identified sequentially from the surface into the interior, by the designations, A, B, C, etc.

Test samples measuring 6 in. (152 mm) long x 1.000 in. (25.4 mm) wide were extracted from the rubber sheets using a die punch. Edge nicks, approximately 0.050 in. (1.27 mm) deep, were introduced using a sharp razor blade. Fractography of samples tested early in this study revealed uneven cut tips. This problem was eliminated with an adjustable centering jig clamped on to the sample while it was being nicked.

2.2 Test Procedure

Samples were tested on an Instron universal testing machine at temperatures ranging from -20 to + 140° C (-4 to + 284° F) using a temperature-controlled chamber that could either be heated or cooled by air circulating through a refrigeration/heating unit. Initially, screw-clamped grips were used with a distance of about 3.5 in. (88.9 mm) between grips. A standard crosshead speed of 0.33 in./s (8.33 mm/s) was used giving a load-displacement strip-chart record. Conversion to strain was based on calibration runs using 1-in. (25.4-mm) gage marks. Later in the program the clamp grips were replaced with Instron elastomeric grips. We also began using an Instron 1-in. gage-length incremental extensometer having 0.10-in. (2.54-mm) increments extending over a 10-in. (254-mm) range. The 0.10-in. extension increments are ticked off on the load-displacement strip-chart record.

In addition to testing at the standard speed, some of the laboratory formulations were tested at stroke-displacement rates of about 235 in./s (6×10^3 mm/s) and 6.6×10^{-4} in./s (1.68×10^{-2} mm/s).

The high stroke rate was obtained on an MTS machine having an open-loop stroke* capability of up to 235 in./s (with a closed-loop stroke capability of up to 15 in./s). The output data of load and extension were fed to a digital transient recording system integrated with a Digital Equipment Corporation LSI-11 minicomputer for data acquisition, processing, and graphics.

A limitation of 10 in. in stroke travel in the high strain-rate system required modifications of the test samples. In addition, in order to reduce the inertia of the sample-and-grip-combination, small aluminum screw-clamped grips were used. For nicked samples, the separation between grips measured 1.63 in. (41.3 mm). Dogbone-shaped strips, with a 0.5 x 0.5 in. (12.7 x 12.7 mm) reduced gage region, were used for the un-nicked samples.

The test records were processed using an HP 9845B desk computer with an HP 9874A digitizer. The results were automatically plotted on an HP 7470A plotter yielding the final stress-strain curves. Both engineering and true stress-strain data were obtained; however, in this paper we present only the engineering values. Stresses reported for nicked samples are nominal stresses, i.e., values are based on the pre-nicked cross sections.

2.3 Evaluation of Tearing Energy

Following the scheme proposed by Oh⁽⁹⁾ the tearing energy, T_t is given by:

$$T_t = 2k(\lambda)cWt \text{ (for thickness } t) \quad (1)$$

where

- $k(\lambda)$ = a proportionality factor varying with the stretch ratio, λ ,
- c = cut depth,
- W = stored strain-energy density far removed from the cut, and
- t = undeformed thickness.

* In the remainder of the report the term crosshead will be used for both crosshead and stroke (ram) displacements.

Dividing by t , we obtain the tearing energy T per unit thickness as:

$$T = 2k(\lambda)cW \quad (2)$$

Using the stress value at which the nicked sample failed as the critical limiting stress on the stress-strain curve of a corresponding un-nicked sample, the value of W is then determined from the area under this curve. A minor error is introduced by considering that all the strain energy is recoverable. The hysteretic loss is assumed to be small compared to variations within a given formulation. The values of c , t , and λ are measured for each sample. The parameter $k(\lambda)$ is determined from the curve given by Oh.⁽⁹⁾

3. RESULTS

3.1 Tearing-Energy Data

The tearing-energy data for the laboratory formulations and commercial materials are plotted in Tables 2 through 8 and 9 through 11, respectively. Under the heading of test numbers the first number refers to the un-nicked sample, the second number to the corresponding nicked samples. The failure stresses refer to the nicked samples. In some cases, the relative tearing energies may appear to be quite inconsistent with the corresponding stress levels. This is due to differences in the sample cut depth, c , used in equation (2). In general, considerable scatter in the data is obtained. However, several trends were detectable.

Figures 1 and 2 contain plots of the average values* of tearing energy obtained at a crosshead rate of 0.33 in./s as a function of temperature for the individual laboratory formulations and commercial

* Average values are used in all plots throughout this report.

pads, respectively. Data for the SBR-35 formulation are included in the plots of Fig. 2. It appears that at this carbon-black content a maximum in tearing energy is obtained at about 60°C. The formulations of SBR-25, -30, and -40 also show either a maximum or at least deviations from the normally expected trend of a decrease in tearing energy with an increase in temperature. The expected trend is obtained with the lower carbon-black contents of SBR-0, -15, and -20. The SBR-35 formulation also shows a maximum at 60°C for the tearing energy results obtained at the crosshead rate of 235 in./s; see Fig. 3. The variation of tearing energy with carbon-black content is shown in Figs. 4 and 5 for crosshead rates of 0.33 and 235 in./s, respectively. At the two lowest temperatures, 22 and -20°C, at the slow strain rate and at all test temperatures at the fast strain rate there is a general increase in tearing energy with an increase in carbon-black content. However, maxima in tearing-energy values are obtained over the range of 25 to 35 wt.% carbon black for samples tested at the three highest temperatures at the slow strain rate.

We obtained very limited data on the effect of strain rate on tearing energy. The results are shown plotted in Fig. 6. Except for one data-point of SBR-35 obtained at the slowest strain rate, the tearing energy increases with an increase in strain rate. An explanation of the various trends and/or anomalies observed was sought in the stress-strain behavior of the corresponding un-nicked samples, in the failure stresses, and in the extent of deviated, knotty, tearing as observed in the failed nicked samples.

3.2 Stress-Strain Curves At 0.33 in./s

Engineering stress-strain curves of un-nicked samples tested to failure at 0.33 in./s at various temperatures are shown in Figs. 7 through 13 for SBR-0, -15, -20, -25, -30, -35, and -40, respectively. Except in a few cases, e.g., for SBR-0, only one curve for each test condition is shown. Note that the insert in Fig. 7 for SBR-0 is drawn on the same scale used for all other formulations.

Except for one abnormally low stress-strain curve at 22°C, the data for SBR-0 are closely grouped together with their positions being random relative to temperature. This also follows with SBR-15. With the higher carbon-black contents, however, the curves obtained at 22° C and/or -20° C are displaced above those curves that are grouped together at the higher temperatures. For all formulations there is a general decrease in failure stress with an increase in temperature.

In Figs. 14 through 18 the stress-strain curves for the un-nicked samples are redrawn to illustrate the influence of carbon-black content at each test temperature. Four main features are illustrated. Firstly, at all temperatures the stress-strain curves are displaced to higher stress values (i.e., an increase in stiffness) with increasing carbon-black content, although some overlapping of the SBR-30 and -35 curves is seen at the three highest temperatures. Secondly, there is a maximum in elongation at about 20 to 25 wt. % carbon black with a sharp fall-off with decreasing amounts of carbon-black content. Thirdly, the failure stress increases with carbon-black content up to about 20 to 25 wt. % and thereafter the variation is random. Fourthly, the slope of the stress-strain curve (i.e., the strain hardening rate) first decreases and then increases with increasing strain. The inflection point, the strain at which a reversal in the change in strain hardening rate occurs, shifts to lower strains with an increase in carbon-black content. However, the inflection point appears to vary randomly with temperature for any given carbon-black content.

Since the only data required from nicked samples for the tearing-energy calculations are the failure stresses, the corresponding stress-strain curves were not processed from the load-displacement records. However, we have shown that for the size of nicks used in this study, the stress-strain curves of the nicked samples virtually superimpose on those obtained from corresponding un-nicked samples, although, depending on the nick depth, the failure stresses may be significantly below those of the un-nicked samples. Also, the trends reported above for the un-nicked samples are consistent with the results reported in a previous study,⁽⁵⁾ which showed the influence of carbon black on stress-strain behavior of single-edge-nicked samples.

3.3 Failure-Stress Plots

Figures 19 and 20 show, respectively, the failure stress as a function of temperature for nicked and un-nicked laboratory-formulated samples. Average values are plotted where more than one sample was tested. Figure 20 shows that for the un-nicked samples the failure stress generally decreases with an increase in test temperature. However, irregular trends are obtained in Fig. 19 for nicked samples that contain more than 20% carbon black. The failure stresses are re-plotted in Figs. 21 and 22 as a function of carbon-black content for the nicked and un-nicked samples, respectively. When plotted in this manner both nicked and un-nicked samples exhibit the same general pattern, namely, a continuous increase in failure stress up to about 25 wt % carbon black followed by one or more maxima. However, exact correspondence in these maxima for a given carbon-black content is not obtained on comparing both sets of plots.

Figures 23 and 24 contain plots of failure stress as a function of strain rate of SBR-0, -15, and -35 for nicked and un-nicked samples, respectively. As may be expected a predominant trend of an increase in failure stress with an increase in strain rate is obtained. However, several deviations from this trend can be noted, especially for SBR-35, which in part could be attributed to the knotty-tearing behavior described below. In addition, some of the samples tested at high strain rates could not be taken to failure due to the test-machine limitations.

3.4 Deviated, Knotty Tearing

In a previous report⁽⁵⁾ we discussed in detail observations on deviated, knotty tearing obtained on deformation of SBR-35; the extent of such behavior increased with either a decrease in strain rate or an increase in temperature. The term "deviated tearing" referred to tearing in a direction that deviates from the normal to the tensile axis, while the term "knotty tearing", as commonly used in the literature,^(2,6,7,8) refers to the knot-like tears, frequently having a

chevron-like character, resulting from the intersection of individual, discontinuous tear paths. In the extreme case, which was not uncommon, the deviation was observed to be 90° , i.e., parallel to the tensile axis. Increases in both strength and tearing energy have been attributed to an effective blunting at the tear tip as the tear deviates from the normal to the tensile loading axis.

For the present paper, we made a cursory examination of the fractured samples listed in Tables 2 through 8. We noted the extent and sharpness of the deviation when it was present. Our observations are summarized in Table 12. The degree of deviated, knotty tearing is rated from 0 to 10, with zero indicating a complete absence of this behavior and 10 a maximum. The ratings are based on the sharpness of the deviation and, to a lesser extent, on the projected length parallel to the tensile axis. We noted two basic types of such tearing. In one type, the entire deviation runs parallel or nearly parallel to the tensile axis and it is terminated with the tearing direction changing abruptly to a direction normal to the tensile axis, becoming unstable and leading to failure. With our samples the extent of this deviation ranges from 0 to a maximum of 3.8 mm projected tear length. Also, the deviated tearing here has markings characteristic of knotty tearing, which can be seen with the naked eye. In the second type, the initiation, progress, and termination of the deviated tearing are all relatively gradual, with the maximum disorientation along a deviation ranging from about 15° to almost 75° from the normal to the tensile axis, with the maximum projected tear length being 5.5 mm. At low magnification ($<10\times$) no markings were observed along these deviated tears. Examples of both types of deviated tearing were illustrated in a previous report.⁽⁵⁾ In Table 12 we list the first type under "sharp" and the second type under "gradual." In some cases a rating is listed under both headings indicating a transition from the first to the second type of behavior. However, in these latter samples, the characteristic knotty tearing markings are either relatively weak or entirely absent. With few exceptions, deviated tearing was limited to the four highest

carbon-black loadings and at the standard and slower strain rates. Some slight indication of deviated tearing is present in SBR-20 samples tested at 140° C and in SBR-35 samples tested at the fast strain rate at 60 to 140° C. In general, the deviated tearing is favored by an increase in temperature and/or a decrease in strain rate; however, a number of exceptions are present. The irregularities in tearing energies and tensile strengths seen in the various plots could be related to the corresponding irregular trends observed in the deviated tearing. This possibility is examined below.

DISCUSSION

The influence of temperature, strain rate, and carbon-black loading on the mechanical properties of SBR formulations has been reported in a number of papers.^(2,7,8,10,11) In general, a decrease in temperature or an increase in strain rate leads to an increase in strength and tearing energy. The addition of carbon black results in large increases in both of these properties as well as developing peak values in the regime where knotty tearing prevails.^(2,8) Our results are generally consistent with these observations.

For our formulations that did not exhibit any deviated, knotty tearing (SBR-0, -15, and -20) the tearing energy increases with either an increase in carbon-black content, a decrease in temperature, or an increase in strain rate (Figs. 1,3-6). The corresponding failure stress for both nicked and un-nicked samples follows the same pattern (Figs. 19-24). Between about 15 and 25 wt.% carbon black large increases in both tearing energy and failure stress occur with an increase in filler content. Between 25 and 40 wt.% carbon black these two properties appear to vary irregularly with the filler content (Figs. 4, 21, 22). The influence of temperature on tearing energy and failure stress for the high carbon-black contents (Figs. 1, 3, 19, 20) only partly follow the trends seen with the low carbon-black contents. There is either a leveling off or a maximum between about 20 to 60° C. The tearing

energies of the commercial pads also follow this general trend; they are generally consistent with the values obtained with the SBR-35 formulation (Fig. 2).

In comparing the engineering stress-strain curves for the different carbon-black contents of unnotched samples tested at a given temperature (Figs. 14-18) several features can be noted:

1. A maximum in failure strain is obtained at some intermediate value of carbon-black content.
2. Over the lower range of carbon black the failure stress increases with an increase in filler content; over the higher range this stress appears to vary randomly at the lower test temperatures, but increases with addition of filler content at the higher test temperatures.
3. The initial stage of deformation exhibits a continuous decrease in strain hardening rate with increasing strain; this is typical of most materials. This behavior is present along the entire curve for all the SBR-0 tests. However, for the carbon-black-loaded formulations an inflection* occurs leading to an increase in the strain hardening rate with strain, virtually until failure. The strain at which the inflection occurs appears to be independent of temperature for a given carbon-black content (Figs. 7-13), but at a constant temperature it appears at increasingly lower strains with an increase in the filler content (Figs. 14-18). There is a slight indication of an inflection point on curves obtained from several SBR-0 samples.

* Although the corresponding true stress-true strain curves are more fundamental and indicative of the instantaneous behavior than are the engineering curves, the inflections we illustrate here are not always detectable on the true curves.

Using the molecular coiled-chain model proposed by Flory,⁽¹²⁾ we can assume that in the early stages of deformation the flow-stress will depend largely upon the relative ease in the uncoiling of the entangled long molecular chains. The flow stress developed during the latter stages of deformation, leading to eventual failure, and the corresponding tearing energy will depend largely upon the resistance to the movement of molecular segments, the breaking of bonds or aggregates of bonds, and the ability to dissipate energy and/or reduce stress concentrations in the vicinity of a tear tip as a tear propagates to final failure. Phenomenologically, the magnitude of these properties is related to the degree of the hysteretic heating developed during the deformation and reflects the viscoelastic behavior of the rubber. Over a wide range of temperatures and strain rates the ability of the long molecular chains, molecular segments and/or crosslinks to respond to the applied loadings decreases with either an increase in strain rate, a decrease in temperature, or an increase in carbon-black loading. At some low temperature (glass transition) or very high strain rate the thermal motions are restricted to the extent that the rubber loses its viscoelastic characteristics, i.e., stress and strain are in phase and the material behaves elastically. It would be expected that the addition of carbon black will facilitate achieving this condition. Similarly, either at some elevated temperature or at a sufficiently low strain rate the thermal motions are able to respond to the applied loadings with the elimination of a phase lag; the material now behaves elastically. Again, one may expect that with carbon-black additions either the temperature will be raised or the strain rate will be lowered at which this condition is achieved.

The exact role of the filler on increasing the inherent resistance of the elastomer to deformation, the initiation of flaw (crack) opening, and the propagation of a crack is still unclear. It is generally considered that a layer of polymer is absorbed onto the surface of the carbon black thereby producing a shell of immobilized elastomer.⁽¹³⁾ This could cause a strengthening not unlike that of particle hardening in crystalline substances. The hysteresis has been attributed to the

breakdown of this structure.⁽¹³⁾ It has also been suggested that the carbon-black particles act as nucleation sites for small tears at highly localized strain regions ahead of the crack, thereby blunting the main tear tip.⁽¹⁴⁾

The critical tearing energy necessary to initiate cut growth in a nicked sample is considered to depend primarily on the molecular structure, and more specifically on the energy necessary to break some minimum number of bonds.^(1,3,8) For example, identical critical values are obtained for both NR and SBR gum vulcanizates. Additions of carbon black can significantly increase these values. Furthermore, the increase in applied energy required for the cut to propagate to some given depth is greater the larger is the hysteresis of the elastomer. This is consistent with the concept that the strengthening by the carbon black is due to some type of modification of the rubber structure that restrains the molecular movements.

One might suggest that the inflection corresponds to a change in the controlling mechanism that resists the applied deformation (strain hardening). Crystallization is certainly an important strengthening mechanism during the straining of natural rubber and this phenomenon may also play a role in SBR. Lyon⁽¹⁵⁾, in a recent study involving calorimetric measurements during tensile straining, noted a significant increase in exothermic behavior starting at the inflection point. It may be that such crystallization take place in SBR.

The details of the processes and forces involved in the uncoiling of the long molecular chains are still unclear. Entropy is certainly a significant factor. One might consider that as the rubber is stretched the role of segmental movements becomes increasingly more important in resisting this stretching. The interactions introduced by the carbon black would result in a general increase in strength. However, the immobilized rubber-shell-carbon-black agglomerates should interfere significantly with the segmental movements. Thus, when the segmental displacements become important for further deformation, an increase in strain hardening rate is initiated (inflection point). The increased

strengthening associated with the larger amounts of carbon black would, in some manner, significantly increase the resistance to the "normal first-stage" uncoiling such that the applied deformation could be more readily accommodated through segmental movements. Thus, with further additions of carbon black the inflection is shifted to increasingly lower strains.

Alternatively, one can consider that the inflection point corresponds to the termination of the uncoiling stage in the rubber phase. Thus, the strain at which the inflection occurs would depend directly upon the relative amount of rubber phase present. In Fig. 25, the inflection strain is plotted versus carbon-black content. The data points represent average values (independent of test temperature); the bars show the range of values. If one considers all ingredients other than the carbon black as part of the rubber phase, then close to a linear relationship is obtained between the inflection strain and rubber contents over the range of 15 to 40 wt.% carbon black (curve A within the range of data points). However, an extrapolation of this near linear behavior to 0 wt.% carbon black (100% rubber phase) would intersect the strain axis at some value considerably below the strains where indications of an inflection point were detected on the SBR-0 stress-strain curves. Inflection points were detectable on several SBR-0 curves at strains ranging from about 155 to 165%. This suggests that the observed values for the inflection strain (Curve A) could be attributed partly to a decrease in the rubber phase (Curve B) and partly to filler strengthening phenomena, per se (Curve C), as the filler content is increased; Curve D refers to the lowest inflection strain that was observed for SBR-0. The concept is illustrated in Fig. 25.

The general decrease in failure strain with an increase in carbon-black content above some intermediate value (≈ 25 wt.%) may be due to dilution of the elastomeric phase by the filler. However, with the exception of the results obtained at 60° C (Fig. 16), the extent of this decrease becomes increasingly greater as the test temperature is lowered from 140 to -20° C. The viscoelastic (hysteretic) behavior should increase with a decrease in temperature over this temperature range. It

has been proposed⁽¹³⁾ that the filler-elastomer interaction decreases with a rise in temperature due to a desorption and the resulting reduction in thickness of the immobilized shells of rubber absorbed on each filler particle. Thus, at the higher temperatures, where the reversal in failure strain is relatively small (Figs. 17 and 18), the filler-dilution effect is, in part, counteracted by this decrease in shell thickness. The decrease in both failure strain and failure stress obtained with an increase in temperature for a given formulation, tested at the standard crosshead speed of 0.33 in./s (Figs. 7-13), places these tests in the lower half of the parabolic-shaped failure envelope⁽⁸⁾. The failure envelope, which gives the failure stress and failure strain at any temperature or temperature-compensated strain rate for a given formulation, shows a maximum failure strain at intermediate temperature/strain-rate values; the strain decreases as the stress either increases in the low temperature/high strain-rate regime or decreases in the high temperature/low strain-rate regime. This behavior is consistent with the expected increased mobility and decreased strength of the elastomer with rising temperature. It may be that the failure strain is a maximum at some intermediate value that corresponds to a maximum in hysteretic (viscoelastic) behavior.

The strength and failure trends observed in our work are consistent with the generally accepted influence of test variables and carbon-black additions on the behavior of SBR formulations. The rapid rate of increase in tearing energy and failure stress between about 15 and 25 wt.% carbon black, especially at the higher test temperatures, coincides with the first appearance of knotty tearing (Figs. 4, 5, 21, 22; Table 12). However, the peaks and valleys along the plotted curves could not always be related to the extent of such knotty tearing. Part of the discrepancies may be due to sample variations; certainly, considerable scatter can be seen in the tabulated data, which list the results of each test sample. However, the regular trends observed for the lower carbon-black formulations give us confidence in the reliability of the data obtained with the higher carbon-black contents. Sampling variations (defects, curing, composition) within a given formulation and minor variations in the nicks may both greatly influence the magnitude

of the deviated, knotty tearing, which, in turn, affects the mechanical properties. In some cases, for a given formulation and test condition, the sample showing the highest resistance to failure exhibits the least amount of knotty tearing. For example, compare in Tables 7 and 12 the SBR-35 tests 136, 153, and 309 all at 140° C. By contrast, the SBR-35 tests 126, 127, and 307 at 60° C exhibit the expected influence of knotty tearing. It is clear from our studies and from studies reported in the literature that maximum or near maximum values of strength and tearing energy are associated with the presence of knotty tearing. However, it is unclear how these maxima are influenced by the depth and geometry of artificial nicks. Based on some cursory studies that we have made on cut depths we believe that this may be an important factor in influencing the degree of knotty tearing and the resulting tearing energy values. As pointed out earlier in this report, precautions were taken to minimize variations in cut depths.

SUMMARY

Tearing energies were obtained on a number of commercial pad materials and on a series of SBR formulations in which the carbon-black content varied from 0 to 40 wt.%. Tests were performed using a standard crosshead rate of 0.33 in./s ($\dot{\epsilon} = 10^{-1}/s$) at 22, 60, 100, and 140° C. The trends obtained with the commercial materials are generally consistent with those exhibited by the formulations having carbon-black contents ranging from 25 to 40 wt.%. The results for the 35-wt.% carbon-black content (close to the carbon-black content of the commercial pads) fall within the range of results obtained for the commercial materials.

For some of the experimental formulations, testing was extended down to -20° C and to both lower ($\approx 2 \times 10^{-4}/s$) and higher ($\approx 470/s$) strain rates. Normal expected trends that could be related to viscoelasticity and carbon-black-rubber- phase interactions were obtained over the range from 0 to 20 wt.% carbon black. Tearing energy and failure stress increased with an increase in filler content and strain rate and decreased with an increase in temperature.

Relatively large increases in tearing energy and failure stress were obtained between about 20 and 25 wt.% carbon black. These increases were greatest at the highest temperature and least at the lowest temperature. The relative increases coincided with the appearance of deviated, knotty tearing.

Between 25 and 40 wt.% carbon black both tearing energy and failure stress generally varied irregularly with either carbon-black content or temperature. Deviated, knotty tearing was exhibited over most of the test variables for these formulations. However, we could not unequivocally relate the observed variations in the data to the tearing behavior. These irregular trends are probably the result of both variations in the sampling and in the razor-blade nicks. Combined, they may cause significant perturbations in the knotty tearing. Failure stress as a function of either carbon-black content or temperature varied in a more systematic manner for the un-nicked samples than for the nicked samples. The phenomenon of deviated, knotty tearing was exhibited to a significantly smaller degree in the un-nicked samples as compared to its presence in nicked samples. In general, a drop in tearing energy and failure stress was obtained with an increase in carbon-black content from 35 to 40 wt.%, and this is attributed to excessive dilution of the rubber phase by the filler. Based on the failure stresses of both nicked and un-nicked samples and on the tearing energies, an optimum carbon-black content between 30 and 35 wt.% is indicated.

Engineering stress-strain curves for un-nicked samples of the experimental formulations are presented and discussed in considerable detail. Maximum elongations were obtained at intermediate values of carbon black. For a given filler content the elongation drops off with an increase in test temperature. All the curves show strain hardening. However, the carbon-black-loaded formulations exhibit an inflection in the change in strain hardening along the stress-strain curve. The inflection appears at increasingly lower strains with increasing carbon-black content; however, for a given carbon-black

content the inflection strain is independent of temperature. The stress-strain behavior and the corresponding trends are discussed in terms of viscoelasticity, rubber networks, carbon-black-rubber-phase interactions, and filler dilution of the rubber phase.

Based on the limited data from the commercial materials together with the data from the 35-wt.% carbon-black laboratory samples, a maximum in tearing energy is developed at about 60° C for these formulations.

RECOMMENDATIONS FOR FURTHER STUDY

1. Examine influence of nick geometry and nick depth on tearing energy relative to sample size for various sample thicknesses. If it is a material property, the critical tearing energy should be constant.
2. Perform a statistical evaluation of the relationship between mechanical properties, (including tearing energy), and the extent and type of knotty tearing of a sufficient number of samples tested under carefully controlled, reproducible conditions.
3. Evaluate the relationship between knotty tearing, type of carbon black, and tearing energy.
4. Clarify the role of carbon black on affecting the inflection strain and strain hardening rates seen on the engineering stress-strain curve.
5. Perform simultaneous calorimetric and strain measurements to ascertain the presence of crystallization and the influence of carbon black on such crystallization in carbon-black-loaded SBR.
6. Expand the study to systematically cover the range of strain

7. Compare the tearing morphologies between commercial materials and experimental formulations and relate these observations to the corresponding tearing energies.
8. Relate tearing energies to hysteretic behavior for different formulations over a matrix of temperatures and strain rates.
9. Evaluate tearing energies of samples previously exposed to cyclic damage simulating field experience.

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AG011

TABLE 1. Designations of Elastomers Tested

A - Laboratory Formulations of Styrene Butadiene Rubber Samples

<u>Ingredient</u>	<u>phr</u>	<u>Ingredient</u>	<u>phr</u>
SBR-1500	100	CBS Santocure	1.0
Sulfur	1.75	Octamine	1.0
Stearic Acid	2.0	Dutrex 726	6.0
Zinc Oxide	5.0	HAF Black-N330	Variable

Carbon-Black (CB)

<u>Designation</u>	<u>phr</u>	<u>wt. %</u>
SBR-0	0	0
SBR-15	20.6	15
SBR-20	29.2	20
SBR-25	38.9	25
SBR-30	50.0	30
SBR-35	62.9	35
SBR-40	77.8	40

Curing: 55 minutes at 145.5° C (294° F); based on t₉₀ for SBR-0 as provided by vendor, Burke Industries, San Jose, CA. Curing time of LLNL compounded materials may have been slightly modified based on t₉₀ obtained by Harold C. Seger, Rubber Consultant.

B - Commercial Materials

<u>Designation</u>	<u>Manufacturer</u>	<u>Track Shoe Design</u>
G-T142	Goodyear	T142
G-T156	Goodyear	T156
F-T142	Firestone	T142
F-T156	Firestone	T156
S-T142	Standard	T142
S-T156	Standard	T156

A number following a designation in Tables 9, 10, and 11 identifies a specific pad, e.g., G-T142-145. In some pads an identification number was not found.

TABLE 2. Tearing Energy Data for SBR-0

<u>Test Numbers</u>	<u>T, °C</u>	<u>Cross-head Rate</u>		<u>Failure Stress</u>		<u>Tearing Energy</u>	
		<u>in./s</u>	<u>mm/s</u>	<u>psi</u>	<u>MPa</u>	<u>in.-lb/in.²</u>	<u>J/m², 10³</u>
225-212	22	236	6x10 ³	197	1358	24	4.20
-213	22	236	6x10 ³	192	1324	26	4.55
618-600	60	236	6x10 ³	96	662	10.1	1.77
-601	60	236	6x10 ³	100	689	10.2	1.79
-621	60	236	6x10 ³	85	586	7.3	1.28
-622	60	236	6x10 ³	113	779	17.4	3.05
623-603	100	236	6x10 ³	78	538	2.0	0.35
624-603	100	236	6x10 ³	78	538	3.6	0.63
626-604	140	236	6x10 ³	75	517	6.2	1.09
-605	140	236	6x10 ³	65	448	4.5	0.79
750-751	22	10	254	153	1055	15.6	2.73
-752	22	10	254	141	972	11.5	2.01
-753	22	10	254	142	979	14.2	2.49
119-121	22	0.33	8.3	90	621	10.2	1.79
-122	22	0.33	8.3	86	593	9.3	1.63
-305	22	0.33	8.3	84	579	8.8	1.54
223-121	22	0.33	8.3	90	621	11.2	1.96
-122	22	0.33	8.3	86	593	9.2	1.61
471-473	22	0.33	8.3	105	724	9.9	1.73
-474	22	0.33	8.3	100	689	11.8	2.07
299-129	60	0.33	8.3	51	352	2.8	0.49
-130	60	0.33	8.3	63	434	6.2	1.09
-306	60	0.33	8.3	65	448	7.3	1.28
293-129	60	0.33	8.3	51	352	3.0	0.53
-130	60	0.33	8.3	63	434	5.8	1.02
141-138	100	0.33	8.3	33	228	1.1	0.19
-139	100	0.33	8.3	49	338	2.6	0.46
-140	100	0.33	8.3	43	296	1.9	0.33
142-143	140	0.33	8.3	41	283	1.3	0.23
-144	140	0.33	8.3	45	310	1.5	0.26
-145	140	0.33	8.3	37	255	1.0	0.18
223-221	22	6.6x10 ⁻⁴	17x10 ⁻³	60	414	11.2	1.96
-222	22	6.6x10 ⁻⁴	17x10 ⁻³	55	379	9.2	1.61

TABLE 3. Tearing Energy Data for SBR-15

Test Numbers	T, °C	Cross-head Rate		Failure Stress		Tearing Energy	
		in./s	mm/s	psi	MPa	in.-lb/in. ²	J/m ² , 10 ³
226-214	22	236	6x10 ³	987	6805	338	59.2
-215	22	236	6x10 ³	860	5930	271	47.5
-216	22	236	6x10 ³	841	5797	269	47.1
619-607	60	236	6x10 ³	217	1496	54	9.46
627-607	60	236	6x10 ³	217	1496	47	8.23
628-608	100	236	6x10 ³	160	1103	40	7.01
-609	100	236	6x10 ³	162	1117	46	8.06
629-605	100	236	6x10 ³	160	1103	31	5.43
-609	100	236	6x10 ³	162	1117	37	6.48
630-610	140	236	6x10 ³	117	807	9.0	1.58
-611	140	236	6x10 ³	144	993	23	4.03
631-610	140	236	6x10 ³	117	807	9.0	1.58
-611	140	236	6x10 ³	144	993	17	2.98
115-116	22	0.33	8.3	443	3054	95	16.6
-117	22	0.33	8.3	307	2117	58	10.2
-118	22	0.33	8.3	280	1931	53	9.28
123-124	60	0.33	8.3	171	1179	33	5.78
-125	60	0.33	8.3	157	1082	28	4.90
-132	60	0.33	8.3	159	1096	29	5.08
135-133	100	0.33	8.3	117	807	10.3	1.80
-134	100	0.33	8.3	109	752	8.4	1.47
-308	100	0.33	8.3	126	869	11.3	1.98
149-150	140	0.33	8.3	99	683	6.1	1.07
-151	140	0.33	8.3	93	641	5.4	0.95
-152	140	0.33	8.3	87	600	4.6	0.81
224-209	22	6.6x10 ⁻⁴	17x10 ⁻³	211	1455	11.3	1.98
-210	22	6.6x10 ⁻⁴	17x10 ⁻³	216	1489	11.6	2.03

TABLE 4. Tearing Energy Data for SBR-20. All Tests Performed at a Cross-head Rate of 0.33 in./s (8.33 mm/s).

<u>Test Numbers</u>	<u>T, °C</u>	<u>Failure Stress</u>		<u>Tearing Energy</u>	
		<u>psi</u>	<u>MPa</u>	<u>in.-lb/in.²</u>	<u>J/m², 10³</u>
370-352	-20	492	3392	92	16.1
-355	-20	471	3247	68	11.9
-358	-20	478	3296	76	13.3
381-352	-20	492	3392	99	17.3
-355	-20	471	3247	75	13.1
-358	-20	478	3296	81	14.2
364-313	22	390	2689	71	12.4
-314	22	389	2682	71	12.4
-325	22	412	2841	78	13.7
328-319	60	331	2282	52	9.11
-331	60	296	2041	48	8.41
-334	60	292	2013	46	8.06
380-319	60	331	2282	51	8.93
-331	60	296	2041	47	8.23
-334	60	292	2013	45	7.88
375-319	60	331	2282	70	12.3
-331	60	296	2041	66	11.6
-334	60	292	2013	62	10.9
361-322	100	250	1724	38	6.66
-337	100	213	1469	26	4.55
-340	100	212	1462	26	4.55
367-343	140	141	972	12	2.10
-346	140	148	1020	15	2.63
-349	140	142	979	12	2.10

TABLE 5. Tearing Energy Data for SBR-25. All Tests Performed at a Cross-head Rate of 0.33 in./s (8.33 mm/s).

<u>Test Numbers</u>	<u>T, °C</u>	<u>Failure Stress</u>		<u>Tearing Energy</u>	
		<u>psi</u>	<u>MPa</u>	<u>in.-lb/in.²</u>	<u>J/m², 10³</u>
371-353	-20	649	4475	105	18.4
-356	-20	634	4371	94	16.5
-359	-20	625	4309	84	14.7
311-315	22	591	4075	102	17.9
-316	22	580	3999	94	16.5
-326	22	569	3923	87	15.2
329-320	60	556	3834	91	15.9
-332	60	683	4709	121	21.2
-335	60	555	3827	91	15.9
362-323	100	557	3840	100	17.5
-338	100	683	4709	104	18.2
-341	100	624	4302	102	17.9
368-344	140	507	3496	81	14.2
347	140	512	3530	88	15.4
350	140	502	3461	79	13.8
376-344	140	507	3496	85	14.9
347	140	512	3530	89	15.6
350	140	502	3461	84	14.7

TABLE 6. Tearing Energy Data for SBR-30. All Tests Performed at a Cross-head Rate of 0.33 in./s (8.33 mm/s).

<u>Test Numbers</u>	<u>T, °C</u>	<u>Failure Stress</u>		<u>Tearing Energy</u>	
		<u>psi</u>	<u>MPa</u>	<u>in.-lb/in.²</u>	<u>J/m², 10³</u>
391-392	-20	690	4757	89	15.6
382-383	22	595	4102	89	15.6
384-385	60	582	4013	86	15.2
386-388	100	660	4551	114	20.0
389-390	140	731	5040	133	23.0

TABLE 7. Tearing Energy Data for SBR-35

Test Numbers	T, °C	Cross-head Rate		Failure Stress		Tearing Energy	
		in./s	mm/s	psi	MPa	in.-lb/in. ²	J/m ² , 10 ³
227-217	22	236	6x10 ³	2241	15,451	383	67.1
-218	22	236	6x10 ³	2091	13,921	369	64.6
228-217	22	236	6x10 ³	2241	15,451	344	60.3
-218	22	236	6x10 ³	2091	13,921	320	56.0
620-612	60	236	6x10 ³	761	5247	574	100.5
-613	60	236	6x10 ³	672	4633	416	72.9
632-612	60	236	6x10 ³	761	5247	551	96.5
-613	60	236	6x10 ³	672	4633	466	81.6
633-614	100	236	6x10 ³	433	2985	326	57.1
-615	100	236	6x10 ³	395	2723	293	51.3
634-614	100	236	6x10 ³	433	2985	164	28.7
-615	100	236	6x10 ³	395	2723	129	22.6
635-616	140	236	6x10 ³	343	2344	66	11.6
-617	140	236	6x10 ³	337	2344	65	11.4
636-616	140	236	6x10 ³	343	2344	98	17.2
-617	140	236	6x10 ³	337	2344	96	16.8
112-108	22	0.33	8.3	719	4957	92	16.1
-109	22	0.33	8.3	826	5695	112	19.6
-110	22	0.33	8.3	659	4544	68	11.9
113	22	0.33	8.3	742	5116	89	15.6
128-126	60	0.33	8.3	921	6350	163	28.5
-127	60	0.33	8.3	687	4737	101	17.7
-307	60	0.33	8.3	871	6005	140	24.5
137-136	100	0.33	8.3	685	4723	83	14.5
-153	100	0.33	8.3	767	5288	118	20.7
-309	100	0.33	8.3	998	6881	158	27.7
148-146	140	0.33	8.3	287	1979	27	4.72
-147	140	0.33	8.3	500	3447	67	11.7
-230	140	0.33	8.3	545	3758	72	12.6
220-211*	22	6.6x10 ⁻⁴	17x10 ⁻³	1558	10,742	292	51.1
-219	22	6.6x10 ⁻⁴	17x10 ⁻³	1000	6895	193	33.8

*Test 211 pulled at cross-head rate of 3.3×10^{-4} in/s.

TABLE 8. Tearing Energy Data for SBR-40. All Tests Performed at a Cross-head Rate of 0.33 in./s (8.33 mm/s).

<u>Test Numbers</u>	<u>T, °C</u>	<u>Failure Stress</u>		<u>Tearing Energy</u>	
		<u>psi</u>	<u>MPa</u>	<u>in.-lb/in.²</u>	<u>J/m², 10³</u>
272-354	-20	739	5095	75	13.1
-357	-20	791	5454	77	13.5
-360	-20	785	5412	76	13.3
312-317	22	768	5295	128	22.4
-318	22	795	5481	131	22.9
-327	22	727	5013	111	19.4
379-317	22	768	5295	92	16.1
-318	22	795	5481	94	16.5
-327	22	727	5013	79	13.8
330-321	60	796	5488	103	18.0
-333	60	684	4716	76	13.3
-336	60	642	4426	76	13.3
363-324	100	583	4020	65	11.4
-339	100	552	3806	55	9.63
-342	100	593	4089	65	11.4
369-345	140	427	2944	45	7.88
-348	140	418	2882	46	8.06
-351	140	461	3179	44	7.88

TABLE 9. Tearing Energy Data for T142 Goodyear Pads-115 and -145 and T156 Goodyear Pad. All Tests Performed at a Crosshead Rate of 0.33 in./s (8.33 mm/s).

<u>Test Number</u>	<u>Layer</u>	<u>T, °C</u>	<u>Failure Stress</u>		<u>Tearing Energy</u>	
			<u>psi</u>	<u>MPa</u>	<u>in.-lb/in.²</u>	<u>J/m², 10³</u>
11	A	22*	950	6550	137	24.0
-7	A	22*	970	6688	142	24.9
-8	E	22*	1030	7102	157	27.5
-9	E	22*	990	6826	150	26.3
64-58	C	60*	1120	7722	177	13.0
-60	D	60*	1090	7515	172	30.1
-61	B	60*	1090	7515	169	29.6
-62	C	60*	1075	7412	163	28.5
70-50	D	100 ⁺	900	6205	147	25.7
-51	D	100 ⁺	665	4585	95	16.6
42	D	100 ⁺	567	3909	70	12.3
71-38	A	140 ⁺	571	3937	87	15.2
-39	A	140 ⁺	380	2620	44	7.71
-40	A	140 ⁺	439	3027	55	9.63
-41	F	140 ⁺	538	3709	79	13.8
25-21	A	22 [#]	892	6150	144	24.7
-22	A	22 [#]	868	5985	132	23.1
-23	C	22 [#]	735	5068	100	17.5
-24	C	22 [#]	735	5068	100	17.5
76-46	B	140 [#]	345	2379	43	7.53
-47	B	140 [#]	391	2696	62	10.9
-48	D	140 [#]	482	3323	71	12.4
-49	D	140 [#]	430	2965	67	11.7

*G-T142-145; ⁺G-T142-115; [#]G-T156.

**TABLE 10. Tearing Energy Data for T142 Firestone Pad-117 and T156 Firestone Pad.
All Tests Performed at a Crosshead Rate of 0.33 in./s (8.33 mm/s).**

<u>Test Number</u>	<u>Layer</u>	<u>T, °C</u>	<u>Failure Stress</u>		<u>Tearing Energy</u>	
			<u>psi</u>	<u>MPa</u>	<u>in.-lb/in.²</u>	<u>J/m², 10³</u>
1-2	A	22*	790	5447	102	17.9
-3	A	22*	880	6067	111	19.4
-4	H	22*	1145	7895	168	29.4
-5	H	22*	1130	7791	153	26.8
65-55	E	60*	983	6778	186	32.6
-57	B	60*	660	4551	79	13.8
-58	C	60*	824	5681	104	18.2
67-53	G	100*	900	6205	129	22.6
-54	G	100*	941	6488	136	23.8
-68	C	100*	687	4737	90	15.8
74-42	D	140*	550	3792	70	12.3
-43	D	140*	555	3827	71	12.4
-44	G	140*	531	3661	66	11.6
-45	D	140*	582	4013	67	13.5
77-88	C	22 ⁺	1330	9170	192	33.6
-89	C	22 ⁺	941	6488	114	20.0
92-103	D	140 ⁺	280	1931	31	5.43
-104	D	140 ⁺	250	1724	20	3.50
-105	D	140 ⁺	495	3413	85	14.9
93-106	B	140 ⁺	555	3827	87	15.2
-107	B	140 ⁺	440	3034	88	10.2

*F-T142-117; ⁺F-T156

TABLE 11. Tearing Energy Data for T156 Standard Pad and T156 Standard Pad-380. All Tests Performed at a Crosshead Rate of 0.33 in./s (8.33 mm/s).

<u>Test Number</u>	<u>Layer</u>	<u>T, °C</u>	<u>Failure Stress</u>		<u>Tearing Energy</u>	
			<u>psi</u>	<u>MPa</u>	<u>in.-lb/in.²</u>	<u>J/m², 10³</u>
81-84	C	22*	1110	7653	183	32.1
-85	C	22*	968	6674	151	26.4
78-86	E	22*	1050	7240	167	29.3
-87	E	22*	1040	7171	162	28.4
94-99	D	140*	600	4137	93	16.3
-100	D	140*	465	3206	65	11.4
77-82	E	22 ⁺	645	4447	87	15.2
-83	E	22 ⁺	616	4247	80	14.0
-90	C	22 ⁺	842	5804	114	20.0
-91	C	22 ⁺	655	4516	83	14.5
95-97	D	140 ⁺	460	3172	67	11.7
-98	D	140 ⁺	360	2482	42	7.36
-101	B	140 ⁺	303	2089	29	5.08
-102	B	140 ⁺	470	3241	51	8.93

*S-T156; ⁺S-T156-380

TABLE 12. Relative Ratings on the Extent of Deviated, Knotty Tearing

Formulation	T, °C	Cross-head		Rating	
		Rate, in./s	Test No.	Sharp	Gradual
SBR-0	all	all	all	0	0
SBR-15	all	all	all	0	0
SBR-20	-20	0.33	all	0	0
SBR-20	22	0.33	all	0	0
SBR-20	60	0.33	all	0	0
SBR-20	100	0.33	all	0	0
SBR-20	140	0.33	343,346	1	
SBR-20	140	0.33	349	2	
SBR-25	-20	0.33	all	0	0
SBR-25	22	0.33	all	0	0
SBR-25	60	0.33	all	2	
SBR-25	100	0.33	323,341	8	
SBR-25	100	0.33	338	9	
SBR-25	140	0.33	344	1	
SBR-25	140	0.33	350	2	
SBR-25	140	0.33	347	3	
SBR-30	-20	0.33	242	0	0
SBR-30	22	0.33	383		2
SBR-30	60	0.33	385	4	
SBR-30	100	0.33	388	2	
SBR-30	140	0.33	390	7	
SBR-35	22	235	all	0	
SBR-35	60	235	612,613		3
SBR-35	100	235	614	0	
SBR-35	100	235	615		2
SBR-35	140	235	616		4
SBR-35	140	235	617		3
SBR-35	22	0.33	108		5
SBR-35	22	0.33	109		4
SBR-35	22	0.33	110		1
SBR-35	22	0.33	113		2
SBR-35	60	0.33	126	8	
SBR-35	60	0.33	127	5	
SBR-35	60	0.33	307	7	
SBR-35	100	0.33	136	8	
SBR-35	100	0.33	153	9	
SBR-35	100	0.33	309	1	
SBR-35	140	0.33	146	10	
SBR-35	140	0.33	147	7	
SBR-35	140	0.33	230	6	
SBR-35	22	3.3×10^{-4}	211	9	
SBR-35	22	6.6×10^{-4}	219	8 1/2	

TABLE 12. Continued

<u>Formulation</u>	<u>T, °C</u>	<u>Cross-head</u> <u>Rate, in./s</u>	<u>Test No.</u>	<u>Rating</u>	
				<u>Sharp</u>	<u>Gradual</u>
SBR-40	-20	0.33	354,318		1
SBR-40	-20	0.33	360		2
SBR-40	22	0.33	317,318		5
SBR-40	22	0.33	327		2
SBR-40	60	0.33	321,336	7	7
SBR-40	60	0.33	333	5	5
SBR-40	100	0.33	324	7	7
SBR-40	100	0.33	342,339	8	8
SBR-40	140	0.33	345,351	8	8
SBR-40	140	0.33	348	9	9

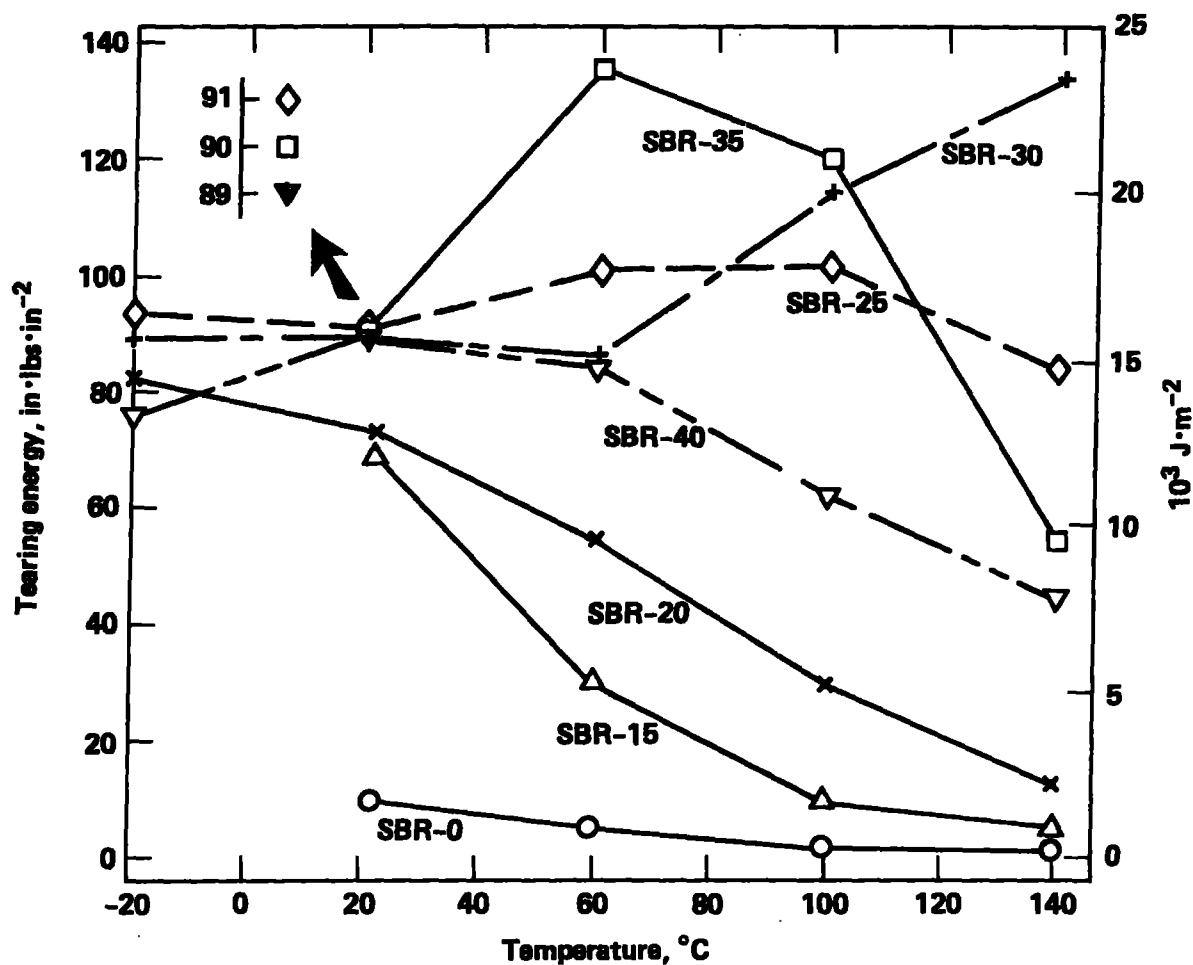


Fig. 1. Tearing energy as a function of temperature for a series of laboratory-formulated SBR elastomers tested at a crosshead rate of 0.33 in./s ($\dot{\epsilon} = 10^{-1}/s$).

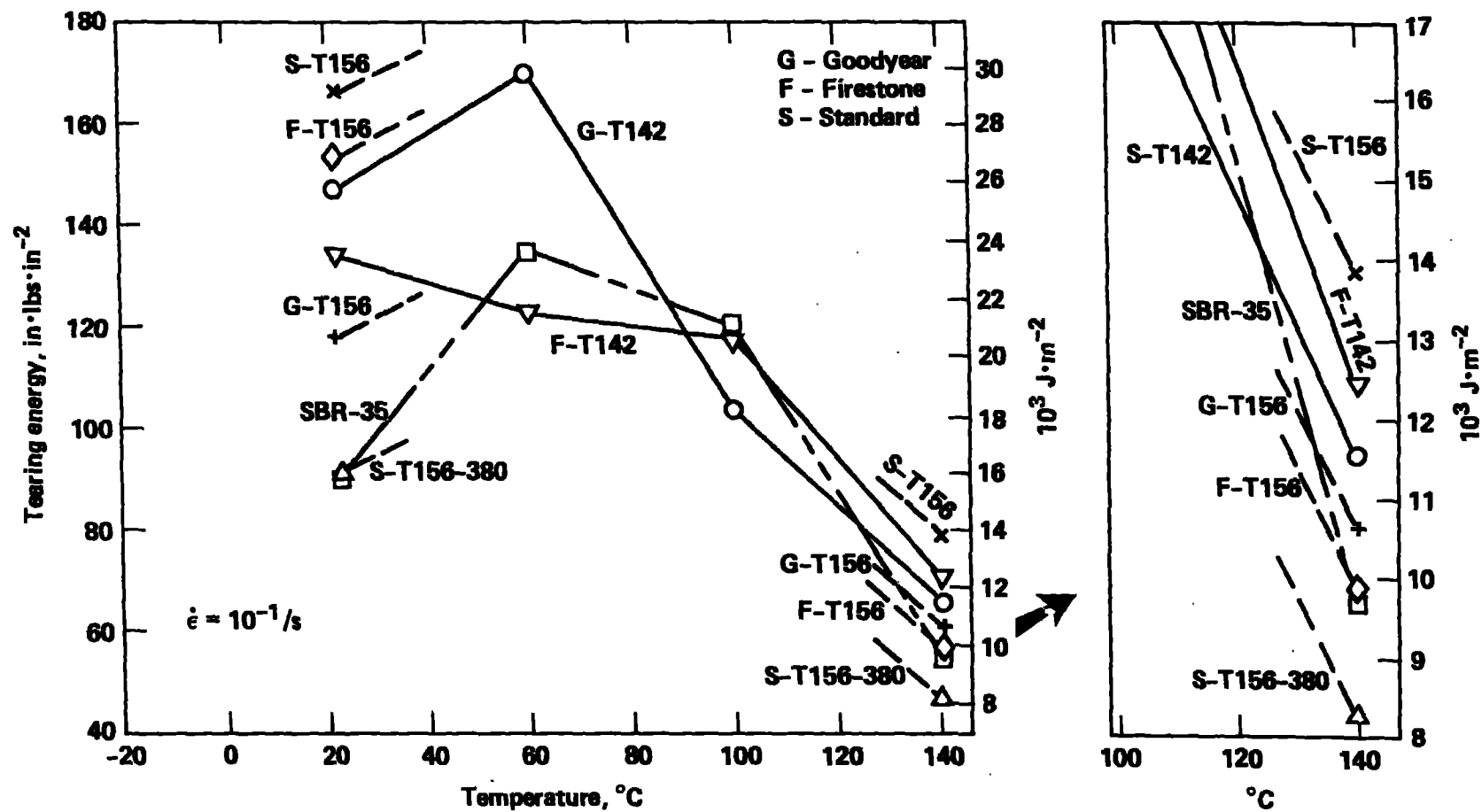


Fig. 2. Tearing energy as a function of temperature for samples removed from tank track pads. Samples tested at a crosshead rate of 0.33 in./s ($\dot{\epsilon} = 10^{-1}/\text{s}$). Results for SBR-35 are included.

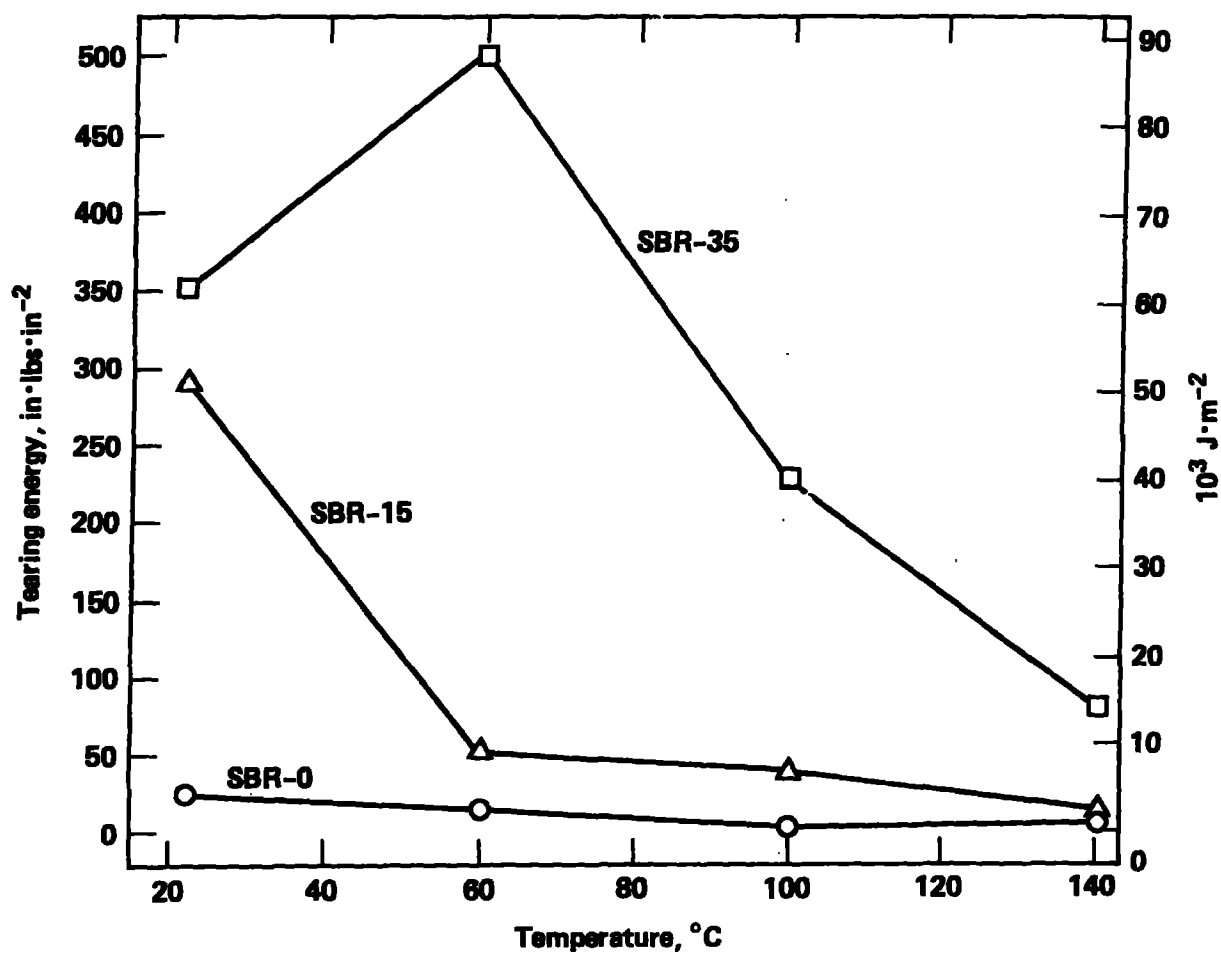


Fig. 3. Tearing energy as a function of temperature for three laboratory-formulated SBR elastomers tested at a high stroke-displacement rate of approx. 235 in./s ($\dot{\epsilon} \approx 470/\text{s}$ for un-nicked samples).

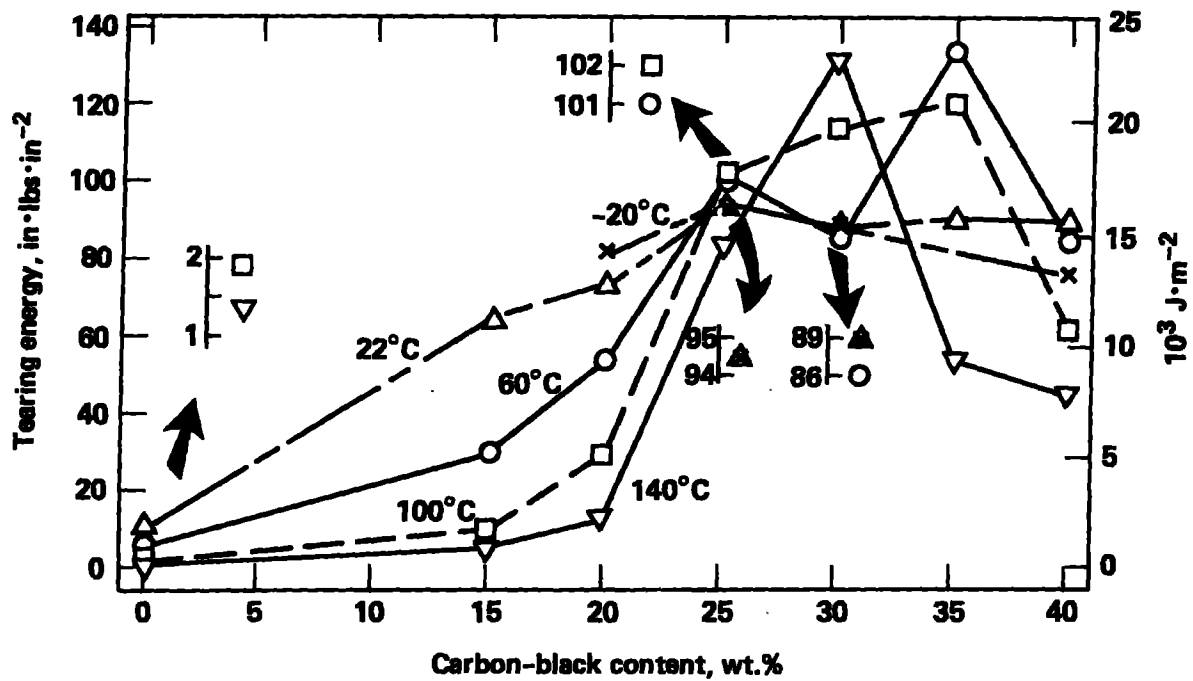


Fig. 4. Tearing energy as a function of carbon-black content. Samples tested at a stroke rate of 0.33 in./s ($\dot{\epsilon} \approx 10^{-1}/\text{s}$) at five temperatures.

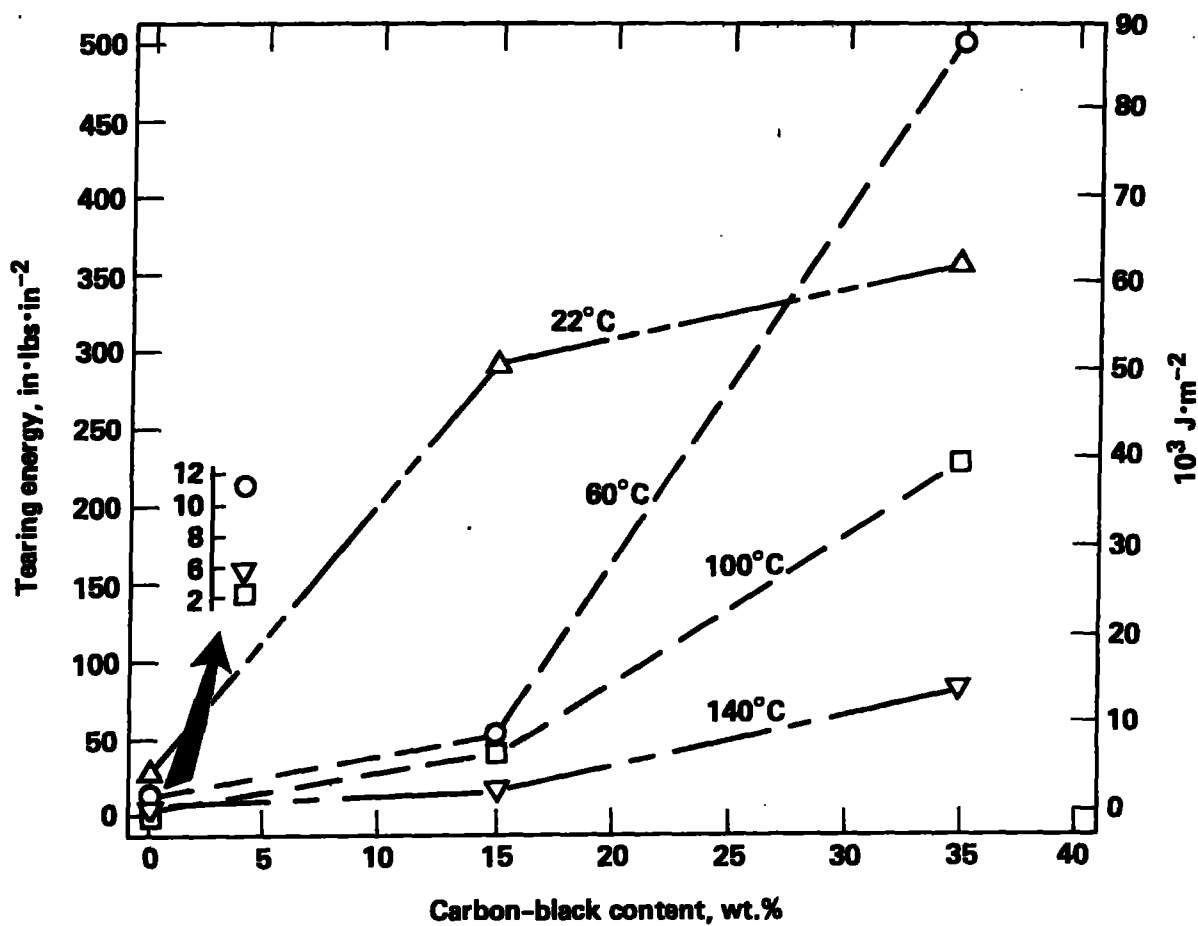


Fig. 5. Tearing energy as a function of carbon-black content. Samples tested at a stroke rate of 235 in./s ($\dot{\epsilon} \approx 470/\text{s}$ for un-nicked samples) at four temperatures.

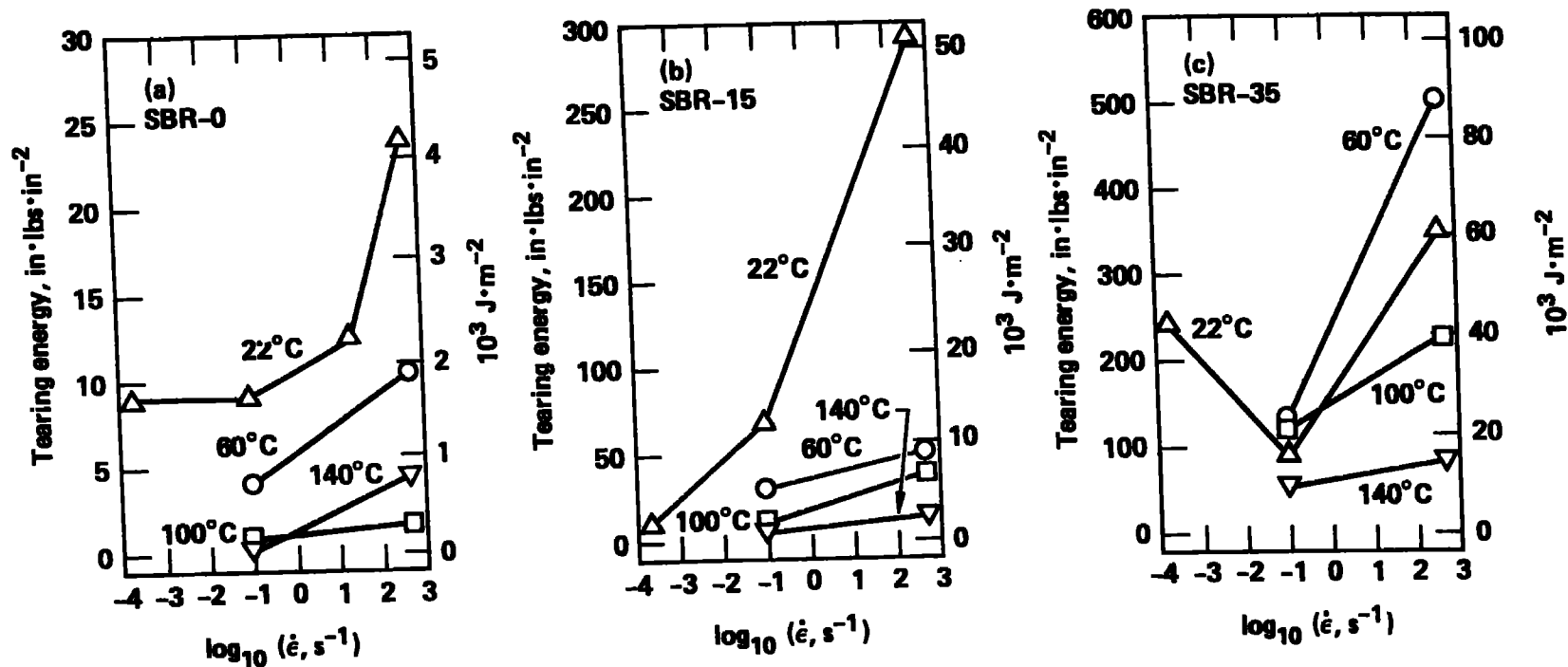


Fig. 6. Effect of strain rate on tearing energy at various temperatures for SBR-0, -15, and -35. (Strain rates are for un-nicked samples).

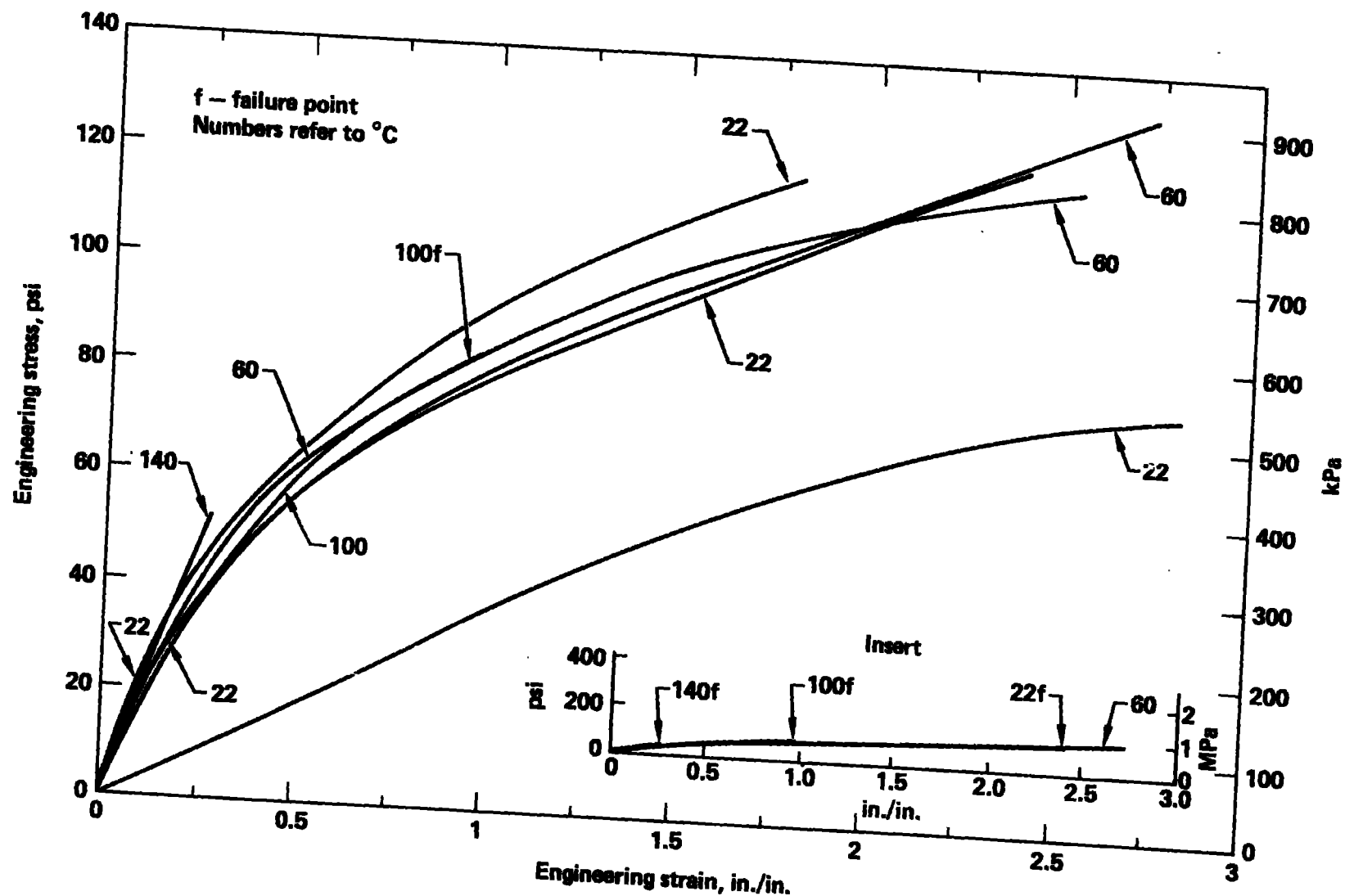


Fig. 7. Stress-strain curves for un-nicked SBR-0 strip samples tested at various temperatures at a crosshead rate of 0.33 in./s ($\dot{\epsilon} \approx 10^{-1}/s$). Insert plotted on same scale used for stress-strain curves of remaining formulations.

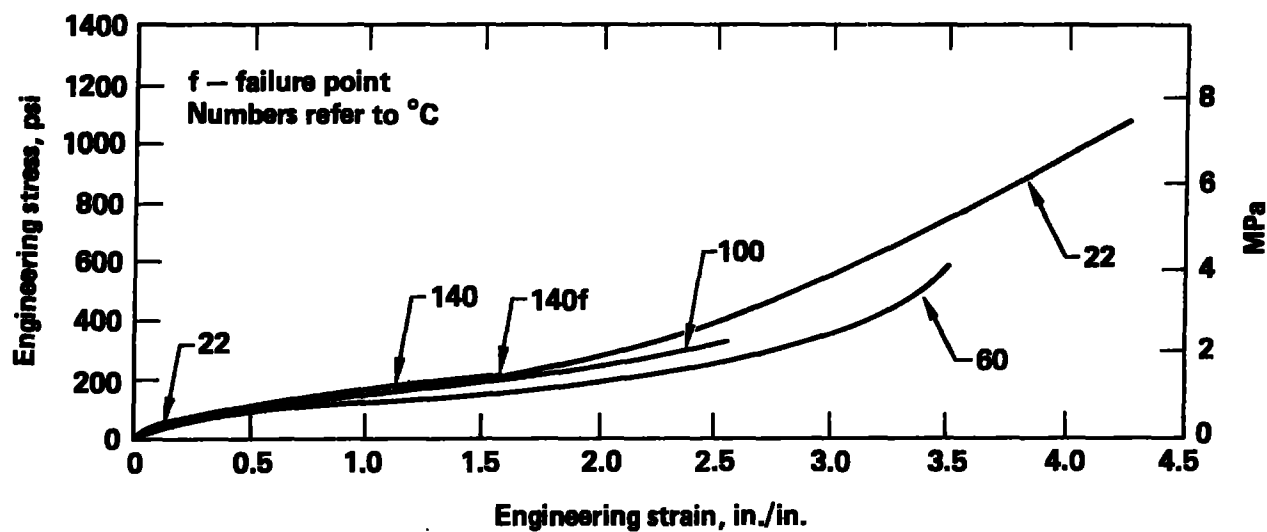


Fig. 8. Stress-strain curves for un-nicked SBR-15 strip samples tested at various temperatures at a crosshead rate of 0.33 in./s ($\dot{\epsilon} \approx 10^{-1}/s$).

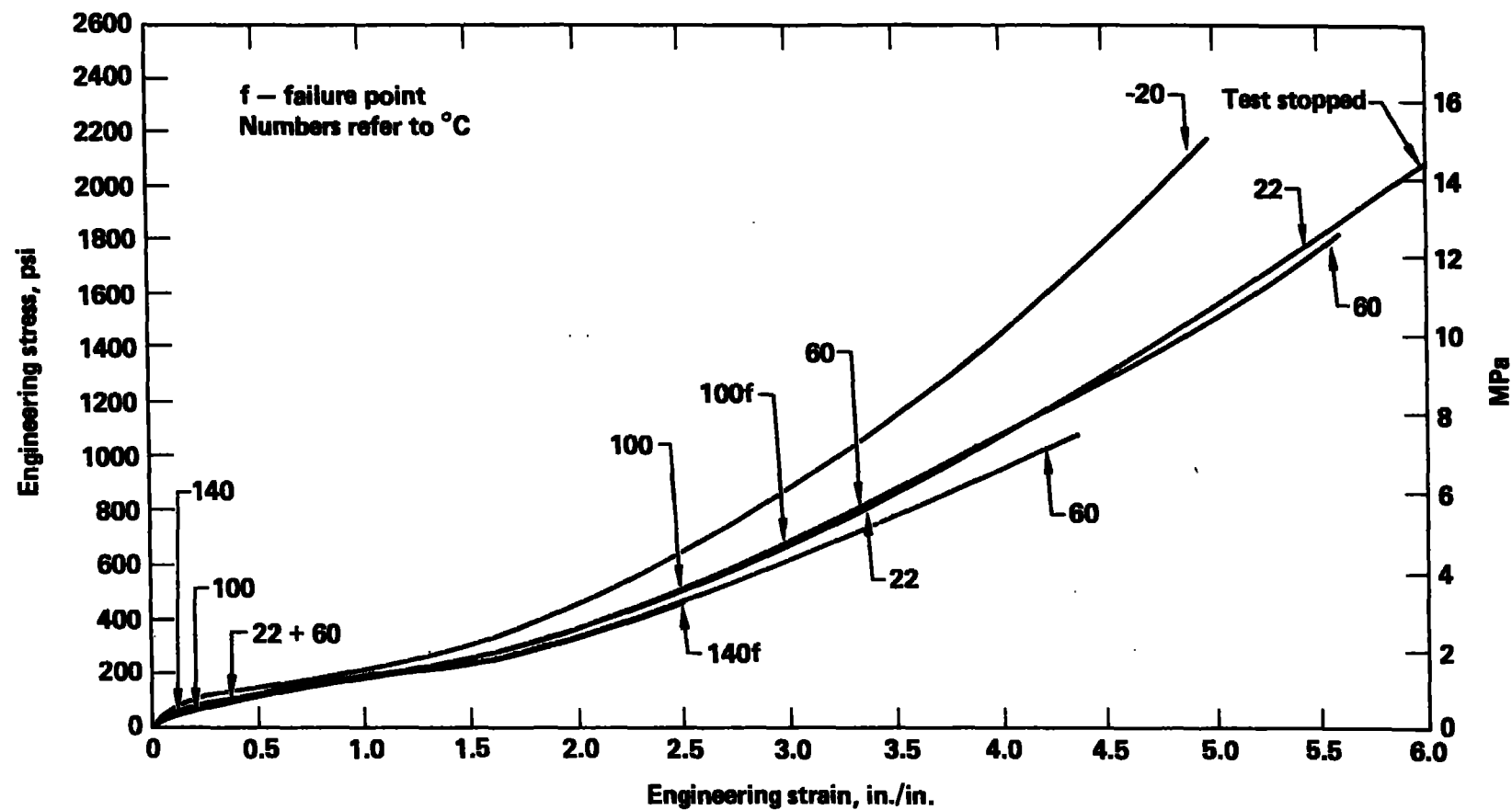


Fig. 9. Stress-strain curves for un-nicked SBR-20 strip samples tested at various temperatures at a crosshead rate of 0.33 in./s ($\dot{\epsilon} = 10^{-1}/s$).

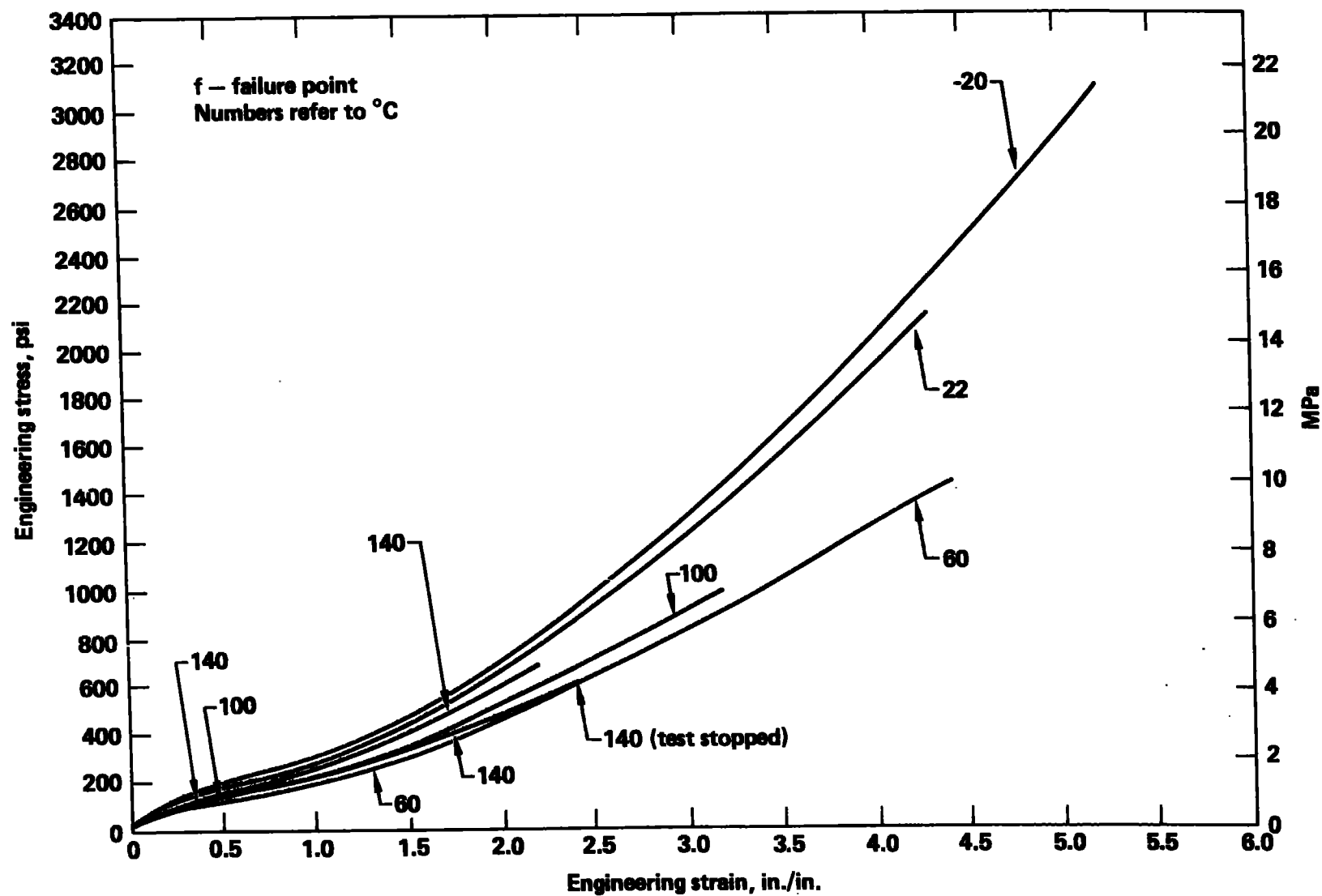


Fig. 10. Stress-strain curves for un-nicked SBR-25 strip samples tested at various temperatures at a crosshead rate of 0.33 in./s. ($\dot{\epsilon} \approx 10^{-1}/s$).

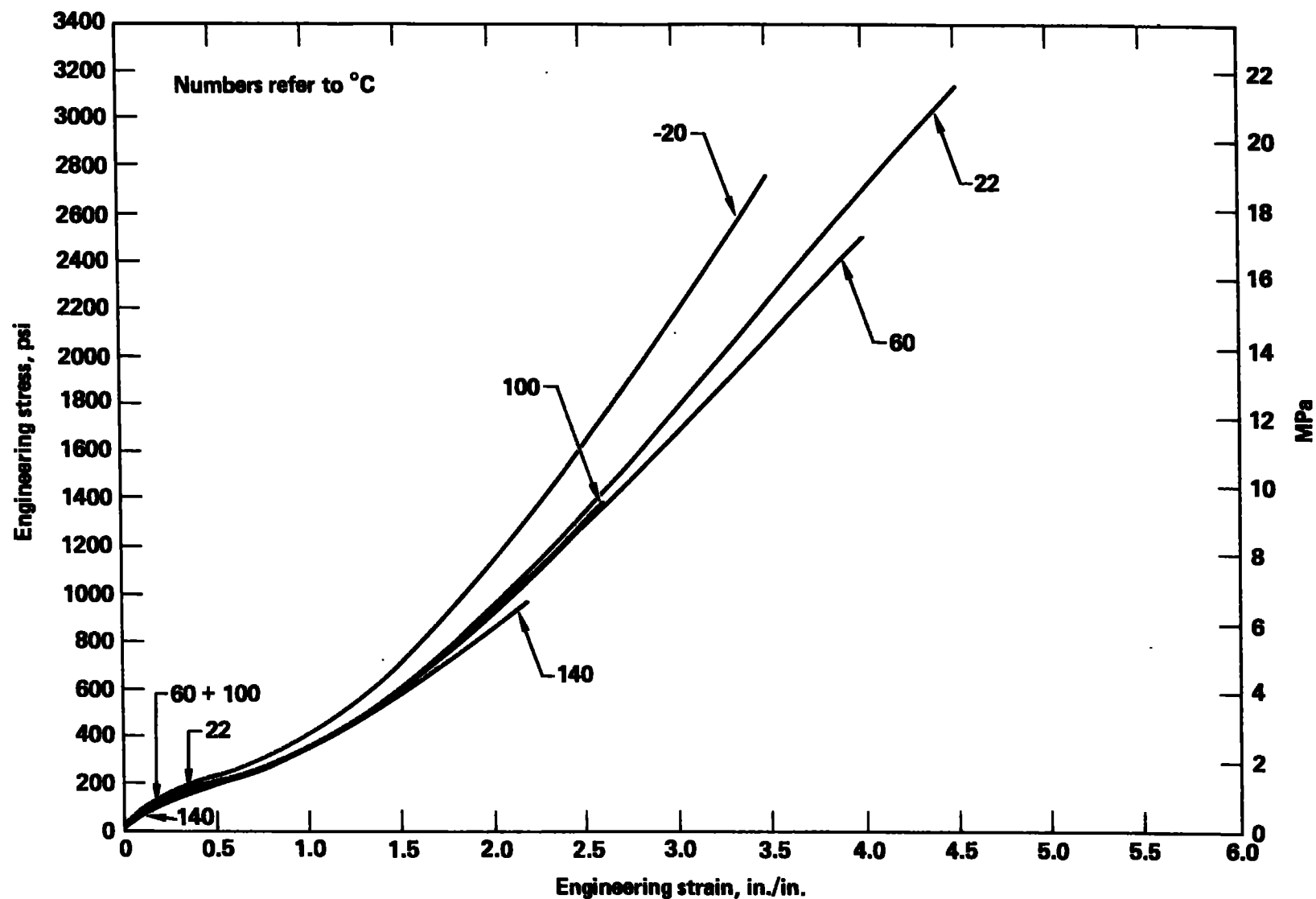


Fig. 11. Stress-strain curves for un-nicked SBR-30 strip samples tested at various temperatures at a crosshead rate of 0.33 in./s ($\dot{\epsilon} \approx 10^{-1}/s$).

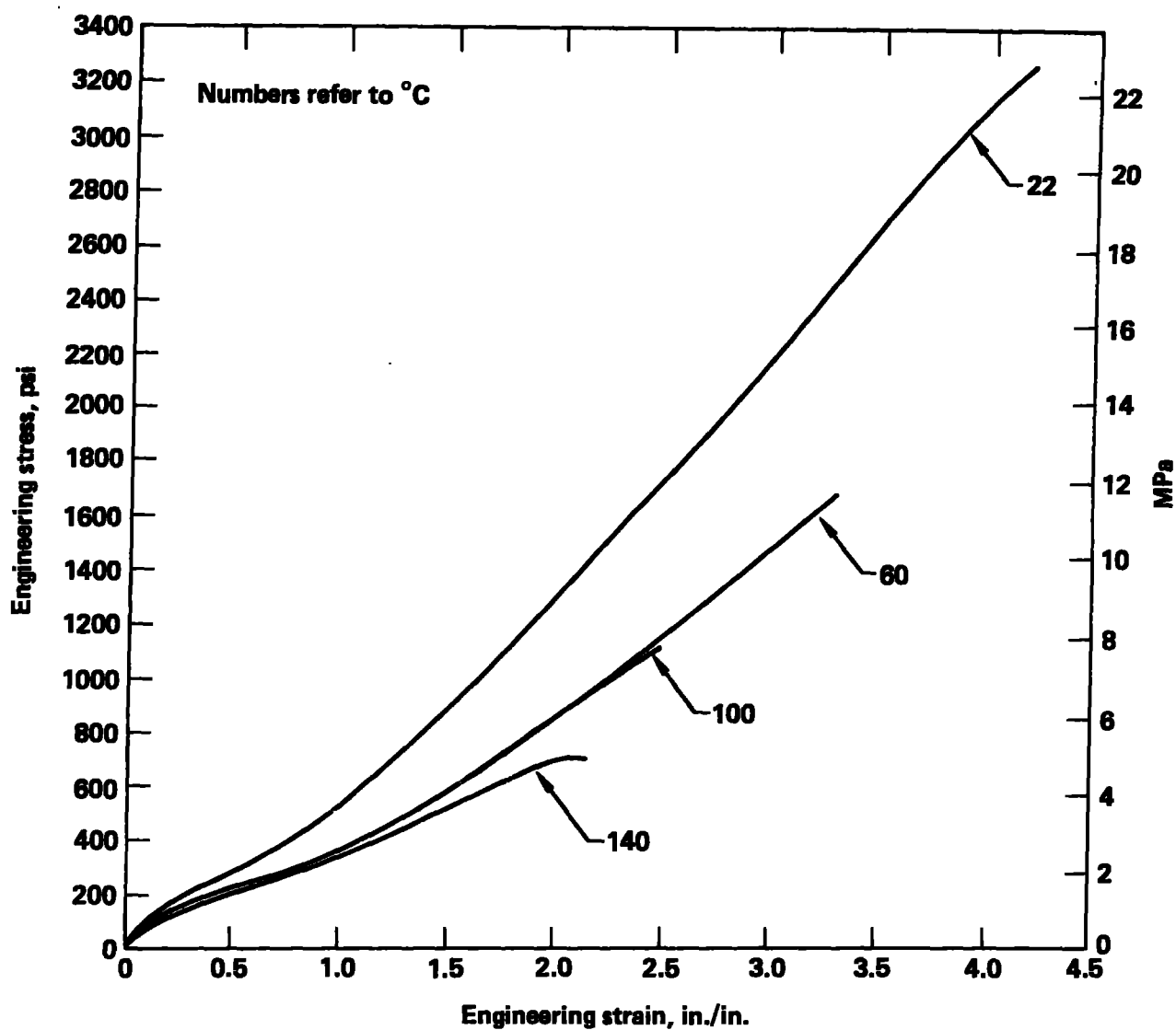


Fig. 12. Stress-strain curves for un-nicked SBR-35 strip samples tested at various temperatures at a crosshead rate of 0.33 in./s ($\dot{\epsilon} \approx 10^{-1}/s$).

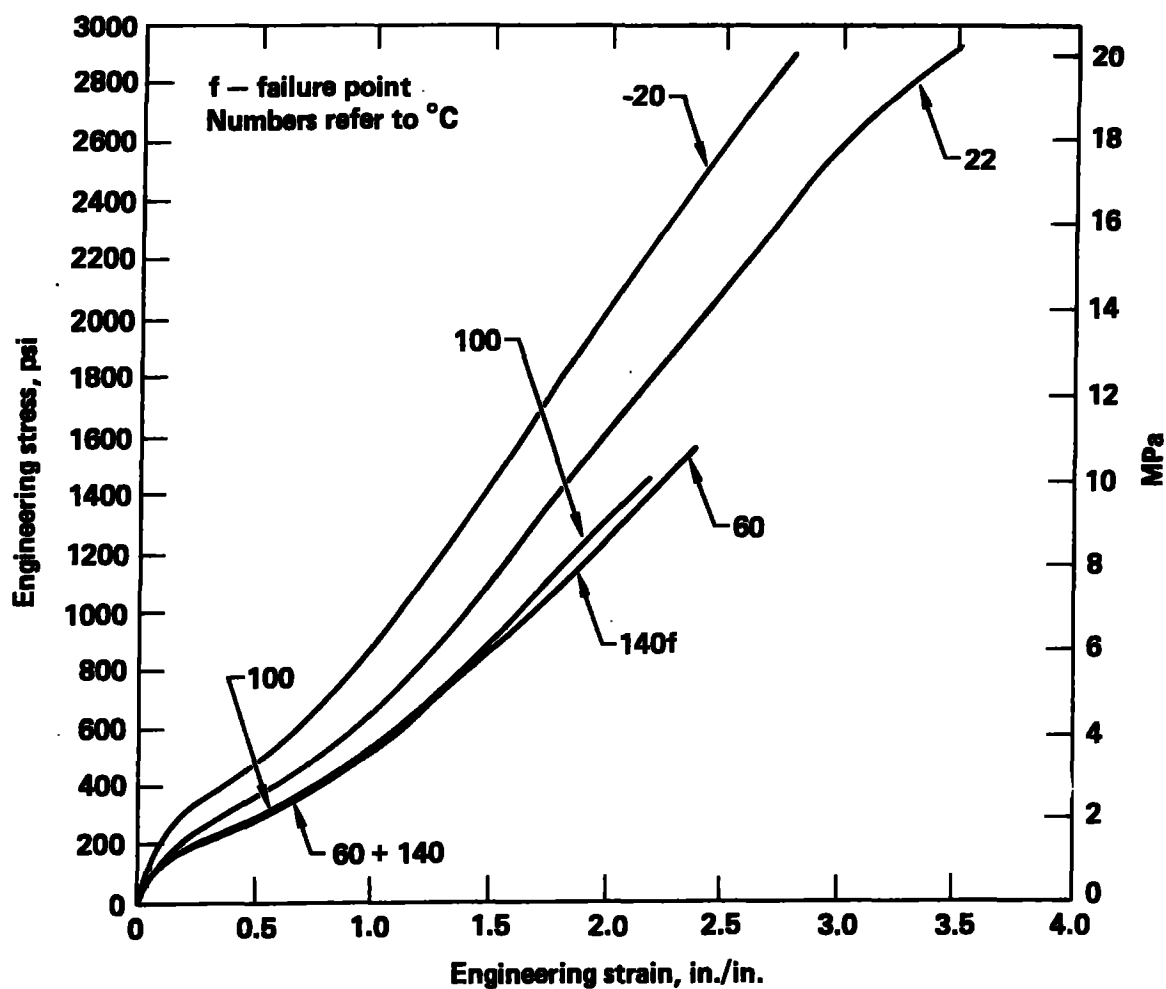


Fig. 13. Stress-strain curves for un-nicked SBR-40 strip samples tested at various temperatures at a crosshead rate of 0.33 in./s ($\dot{\epsilon} \approx 10^{-1}/s$).

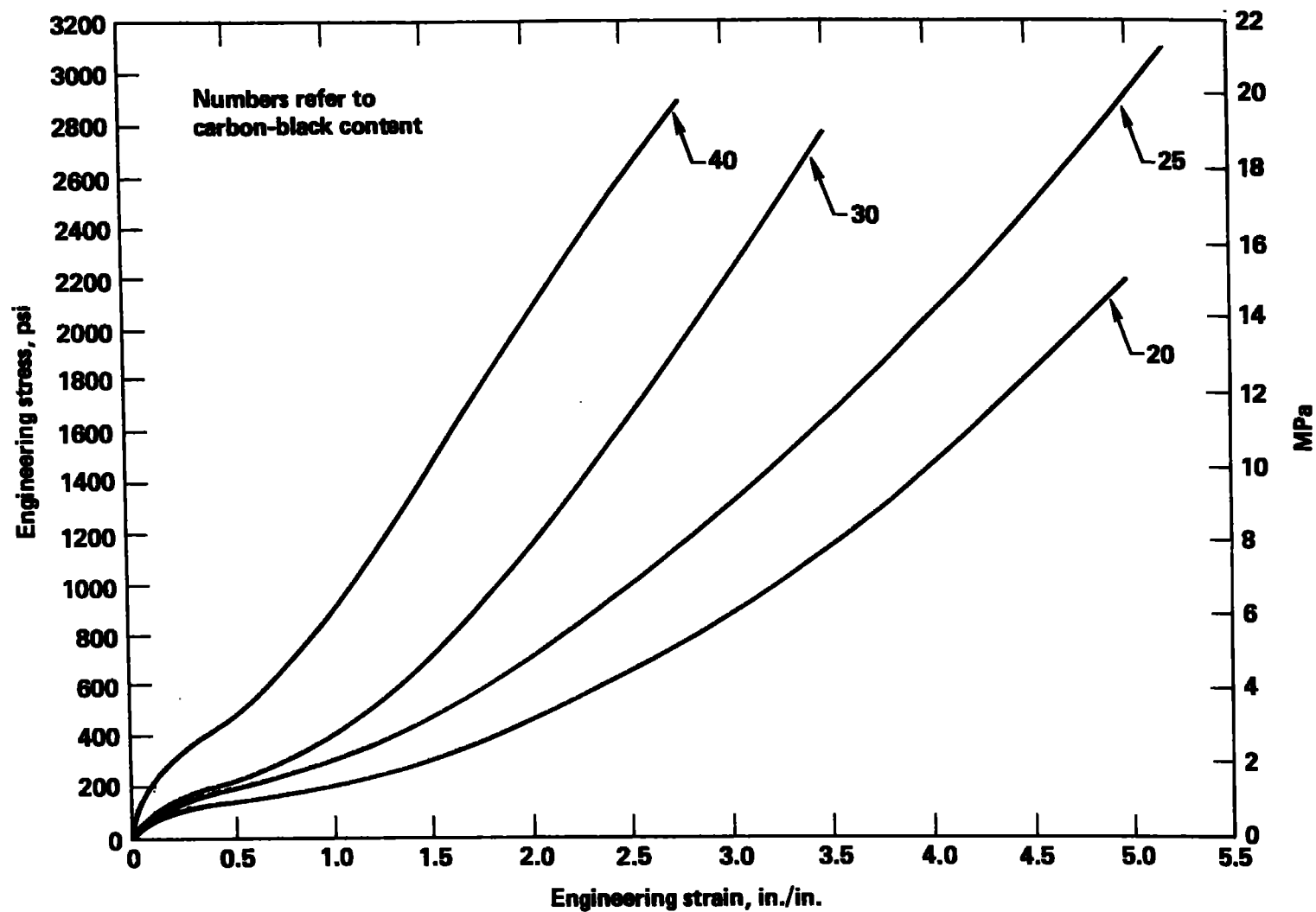


Fig. 14. Stress-strain curves for un-nicked SBR strip samples tested at -20°C at a crosshead rate of 0.33 in./s ($\dot{\epsilon} \approx 10^{-1}/\text{s}$).

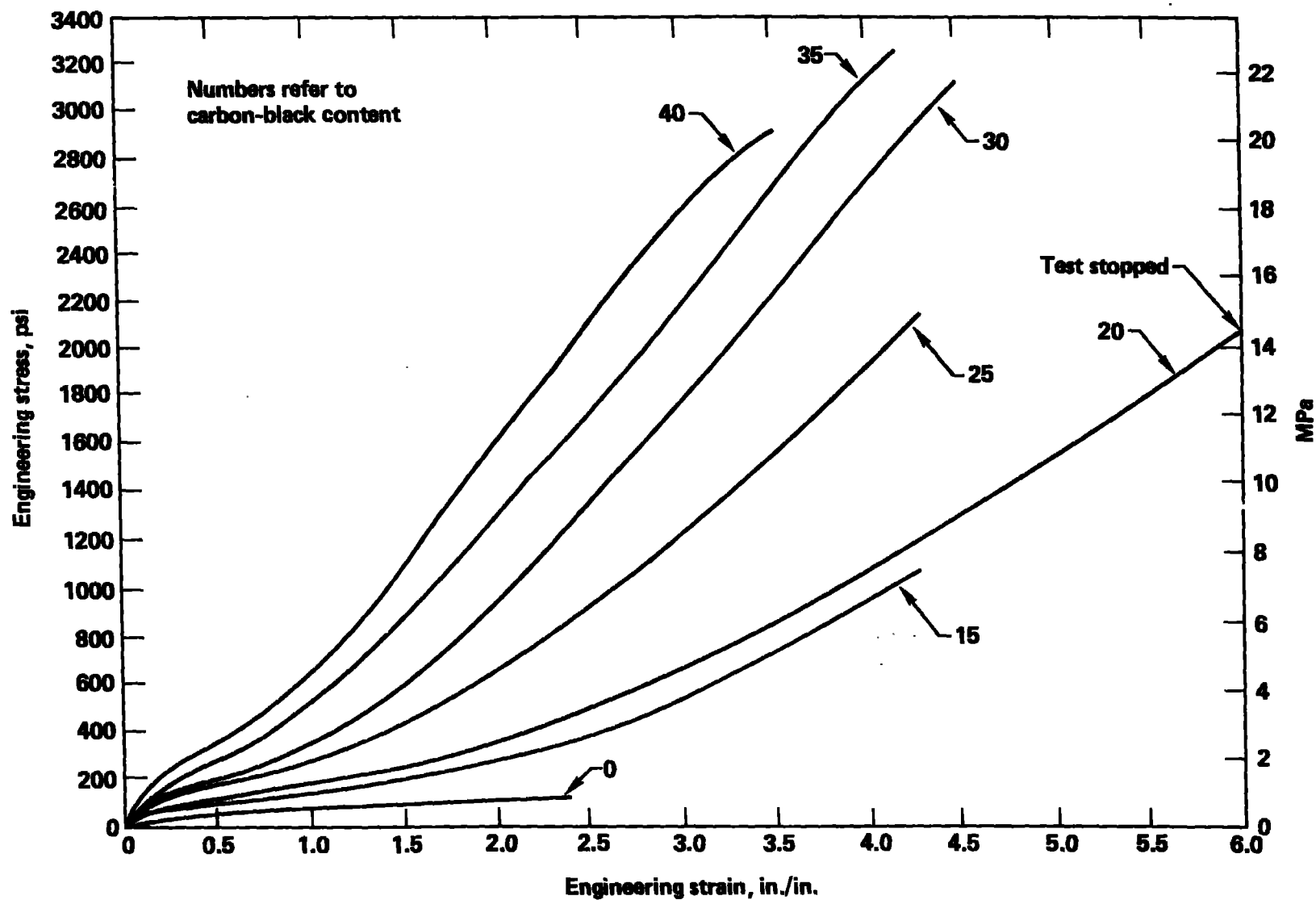


Fig. 15. Stress strain curves for un-nicked SBR strip samples tested at 22° C at a crosshead rate of 0.33 in./s ($\dot{\epsilon} = 10^{-1}/s$).

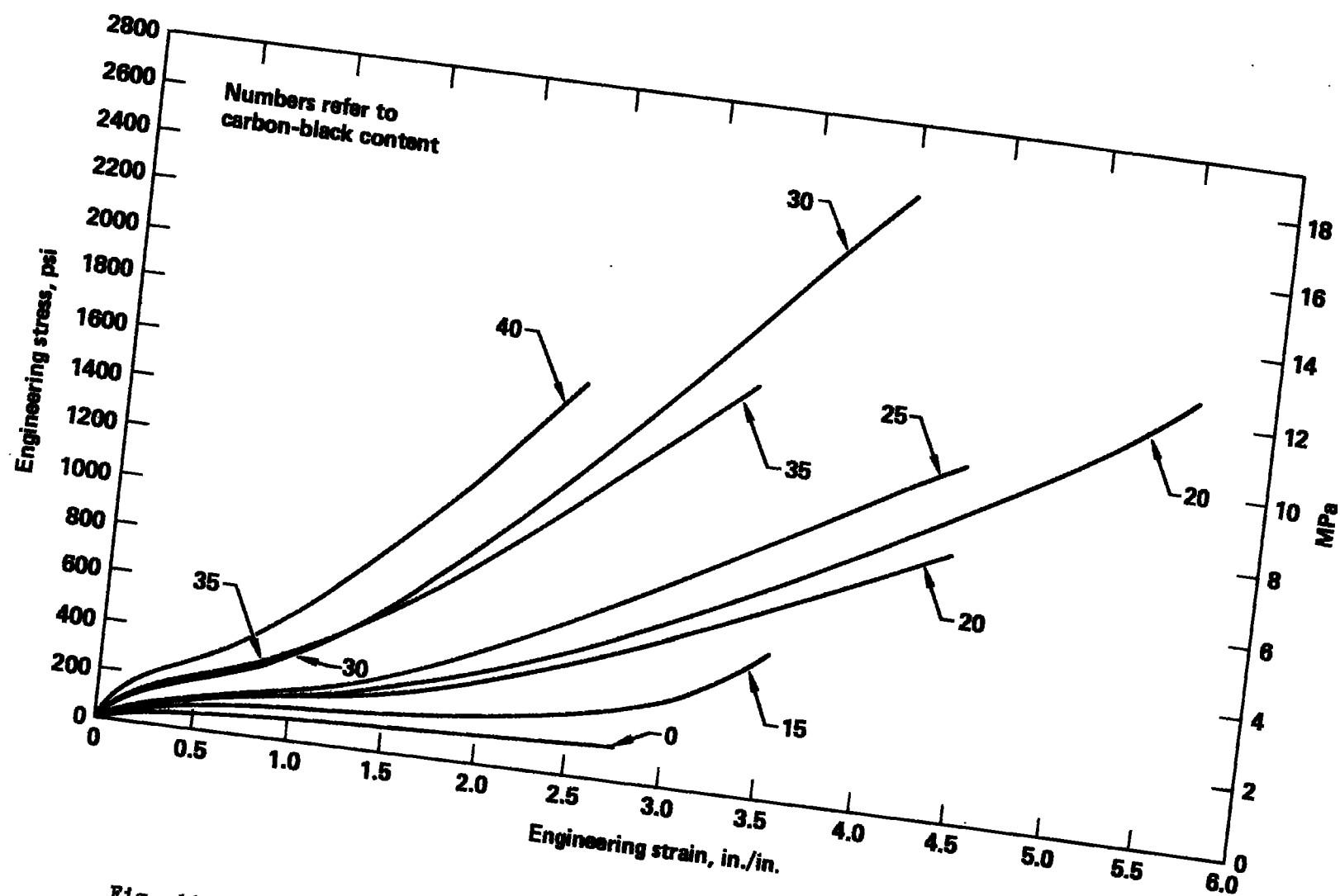


Fig. 16. Stress-strain curves for un-nicked SBR strip samples tested at 60° C at a crosshead rate of 0.33 in./s ($\dot{\epsilon} \approx 10^{-1}/s$).

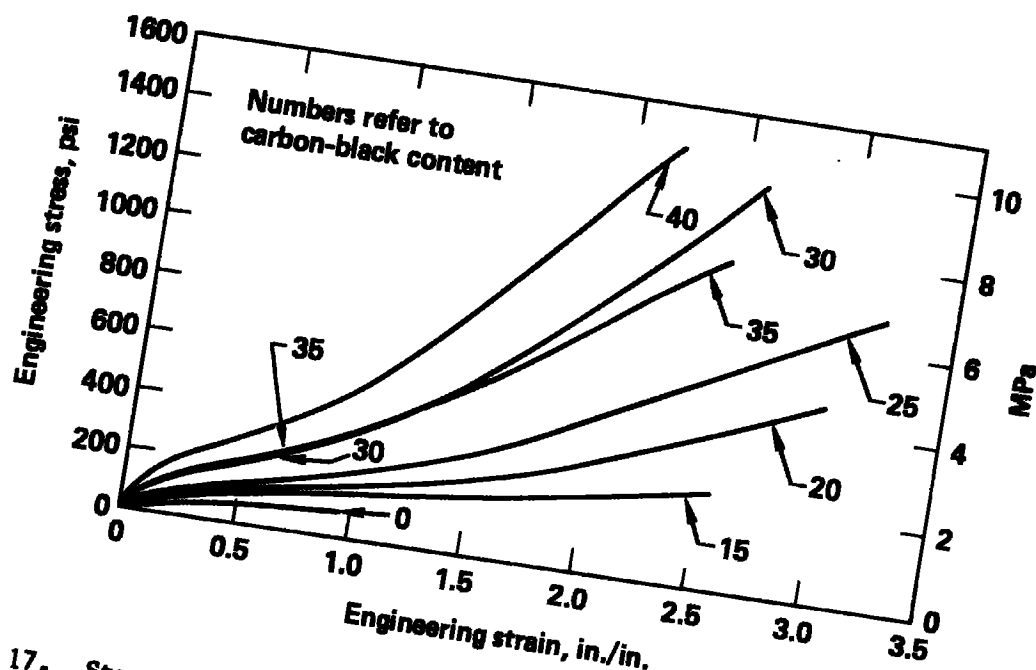


Fig. 17. Stress-strain curves for un-nicked SBR strip samples tested at 100° C at a crosshead rate of 0.33 in./s ($\dot{\epsilon} \approx 10^{-1}/s$).

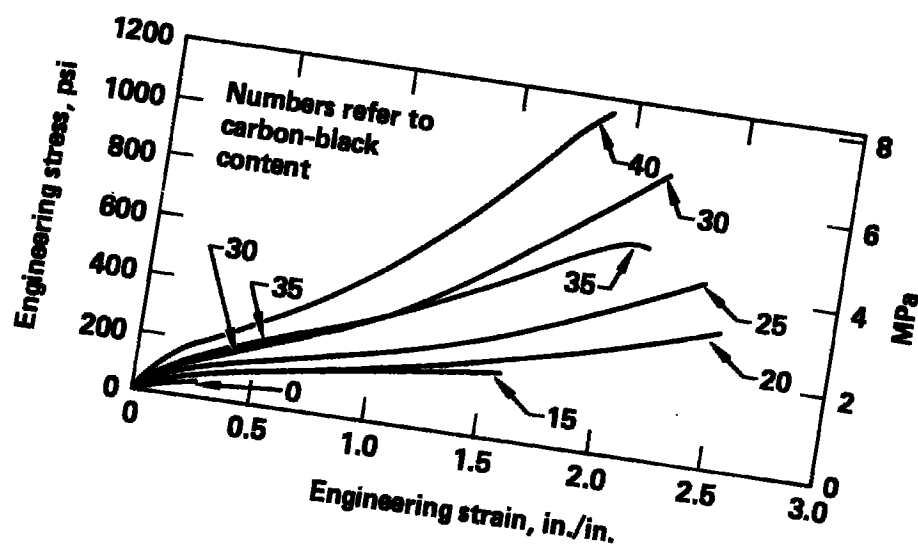


Fig. 18. Stress-strain curves for un-nicked SBR strip samples tested at 140° C at a crosshead rate of 0.33 in./s ($\dot{\epsilon} \approx 10^{-1}/s$).

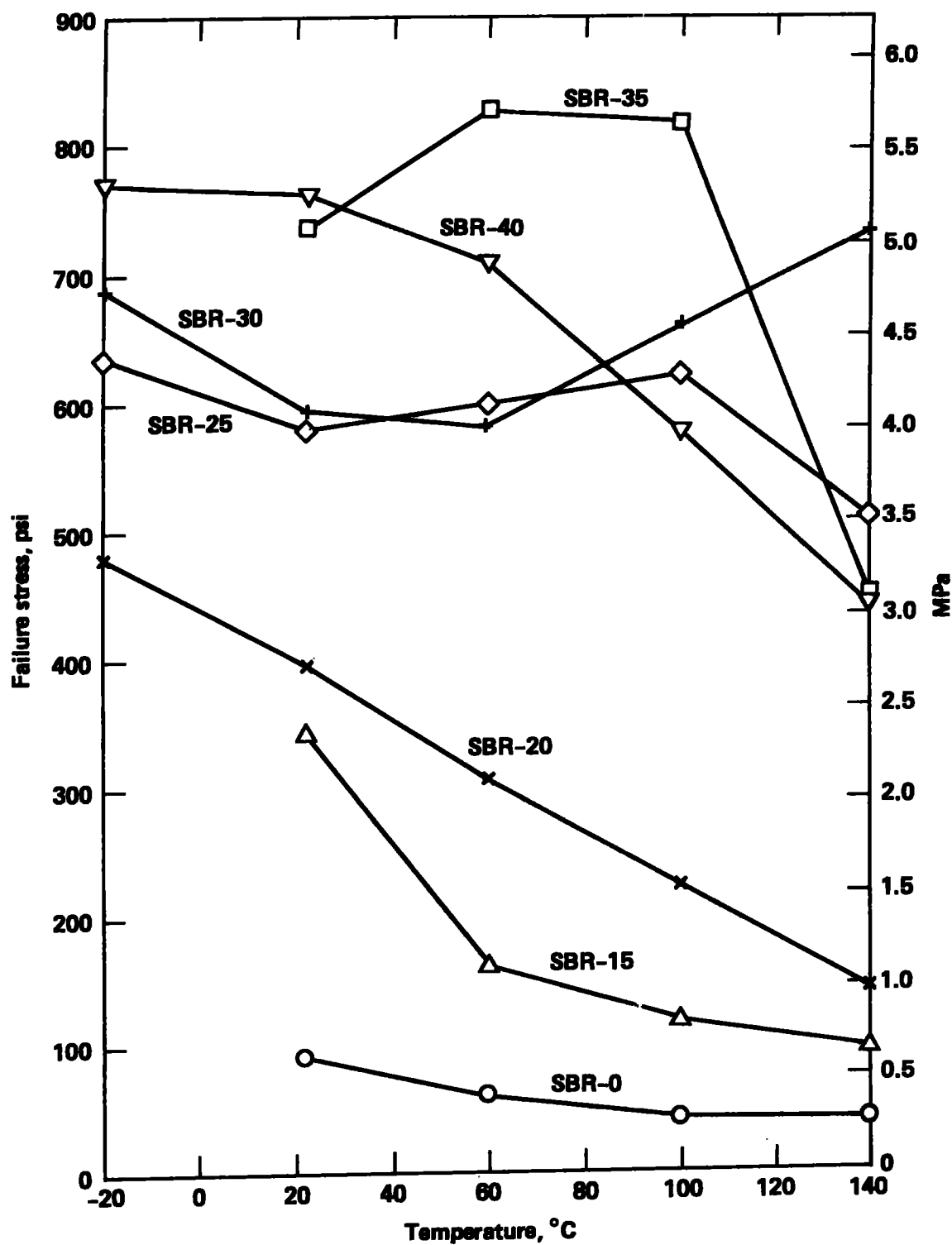


Fig. 19. Failure stress of nicked SBR strip samples tested at various temperatures at a crosshead rate of 0.33 in./s ($\dot{\epsilon} \approx 10^{-1}/s$).

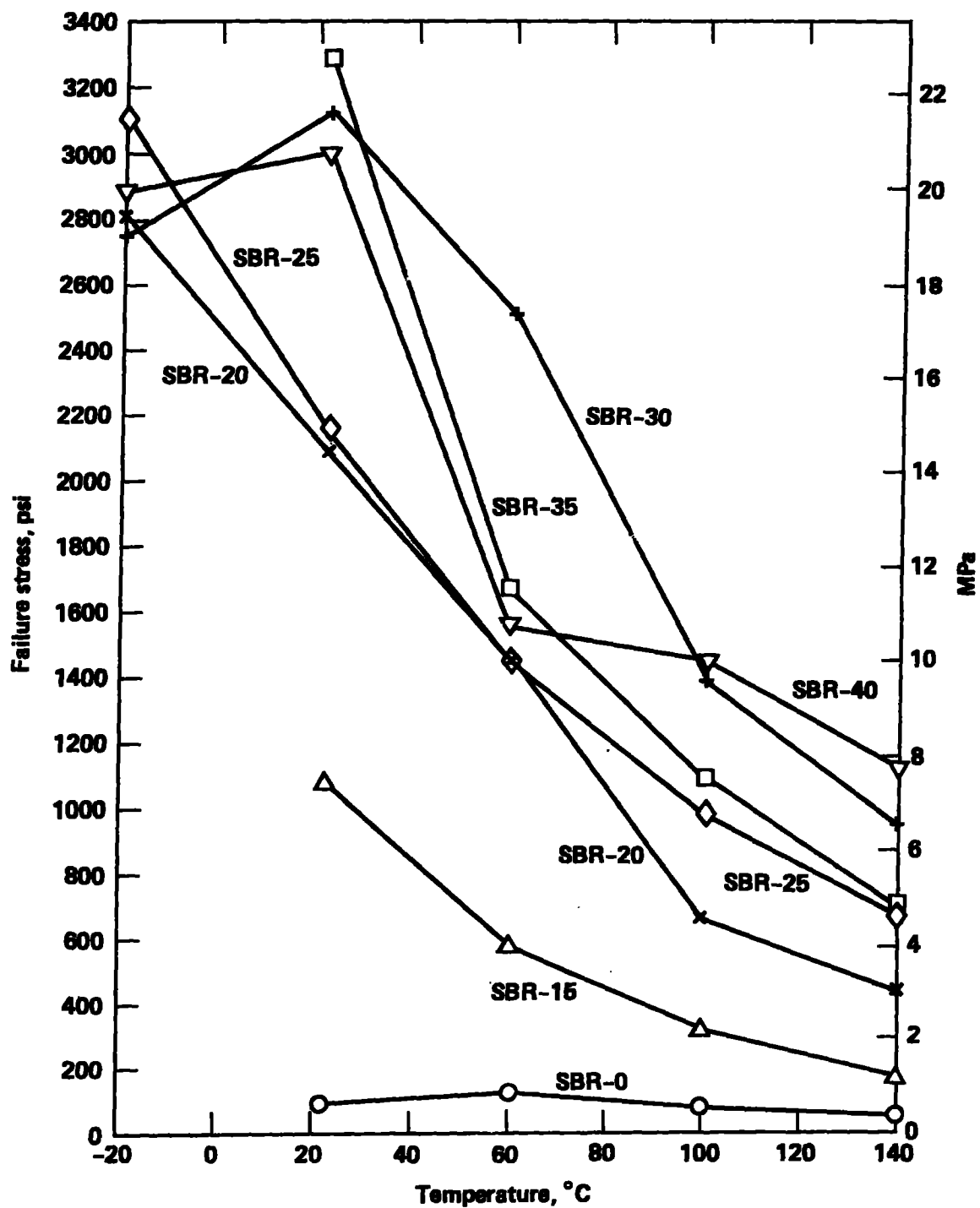


Fig. 20. Failure stress of un-nicked SBR strip samples tested at various temperatures at a crosshead rate of 0.33 in./s ($\dot{\epsilon} \approx 10^{-1}/s$).

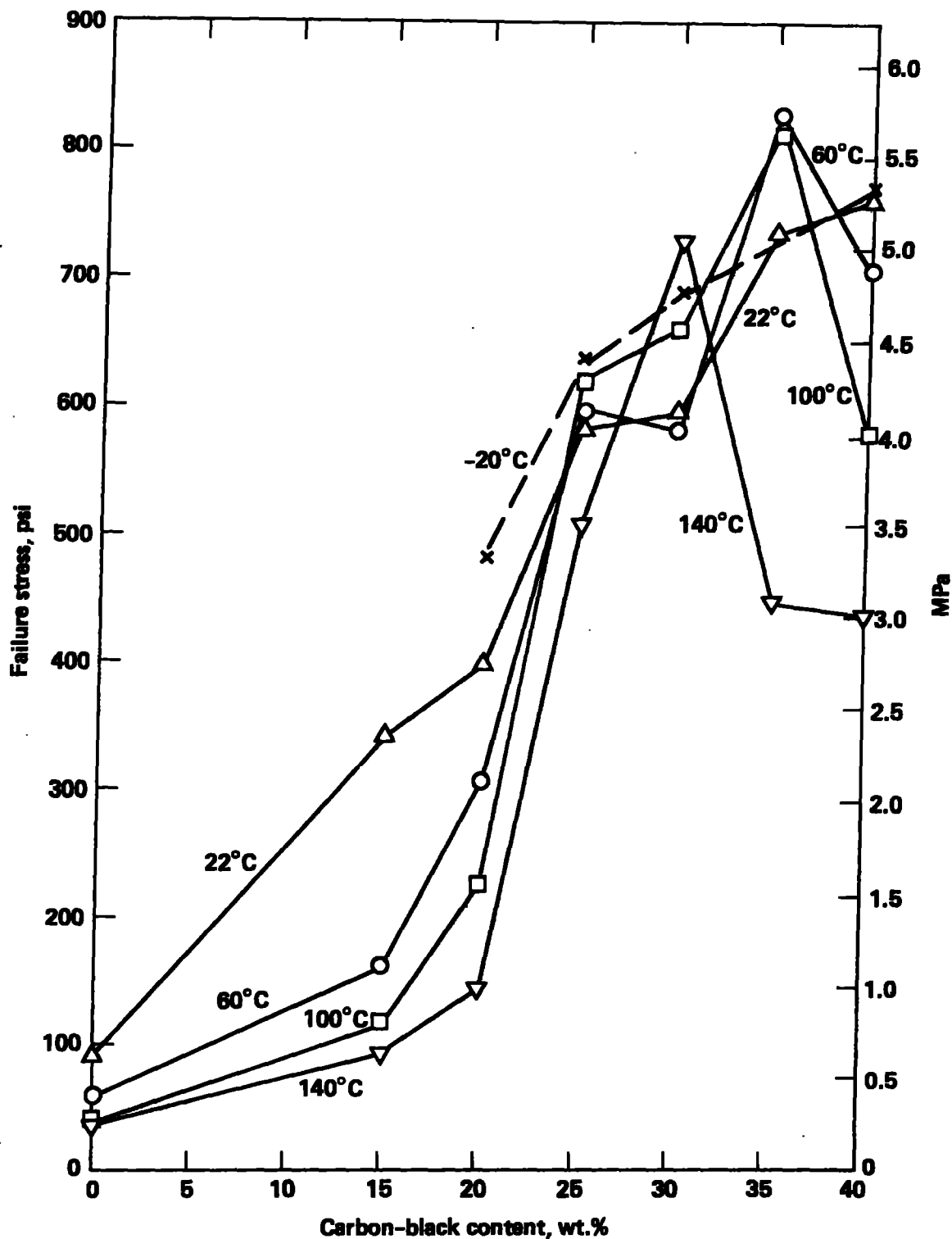


Fig. 21. Failure stress of nicked SBR strip samples as a function of carbon-black content at a series of temperatures. Tested at a crosshead rate of 0.33 in./s ($\dot{\epsilon} \approx 10^{-1}/s$).

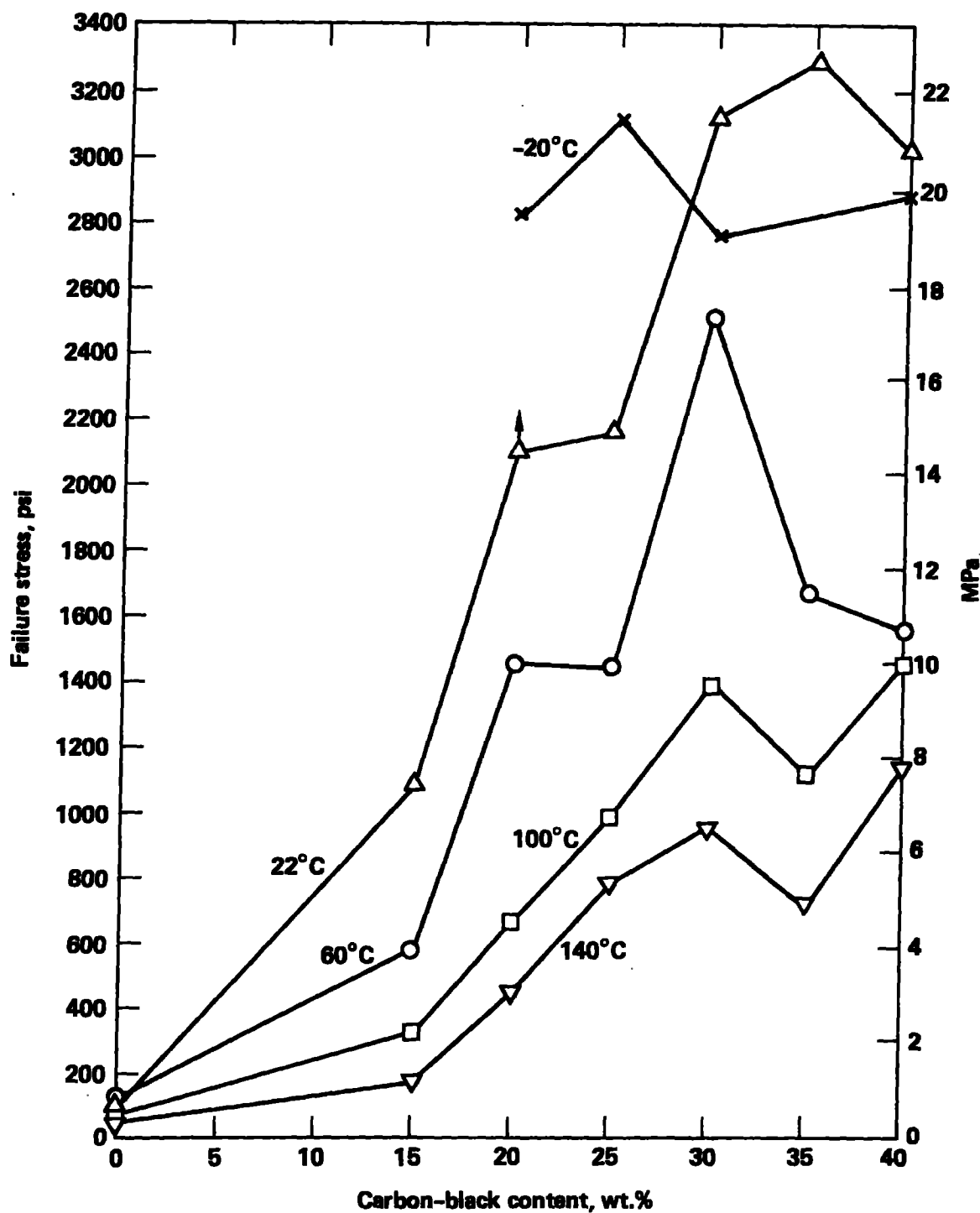


Fig. 22. Failure stress of un-nicked SBR strip samples as a function of carbon-black content at a series of temperatures. Tested at a crosshead rate of 0.33 in./s ($\dot{\epsilon} \approx 10^{-1}/s$).

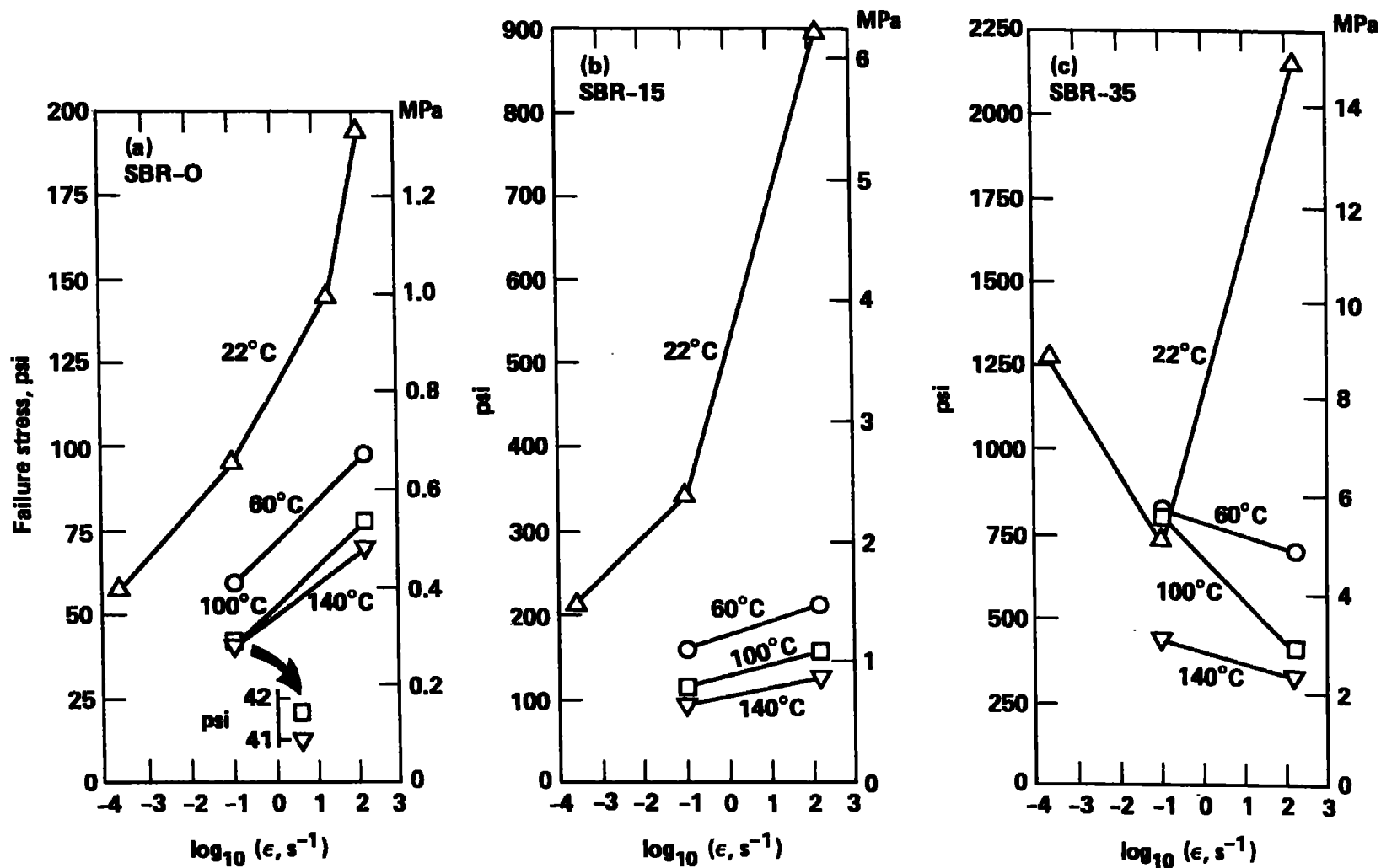


Fig. 23. Effect of strain rate on failure stress at various temperatures for nicked strip samples of SBR-0, -15, and -35.

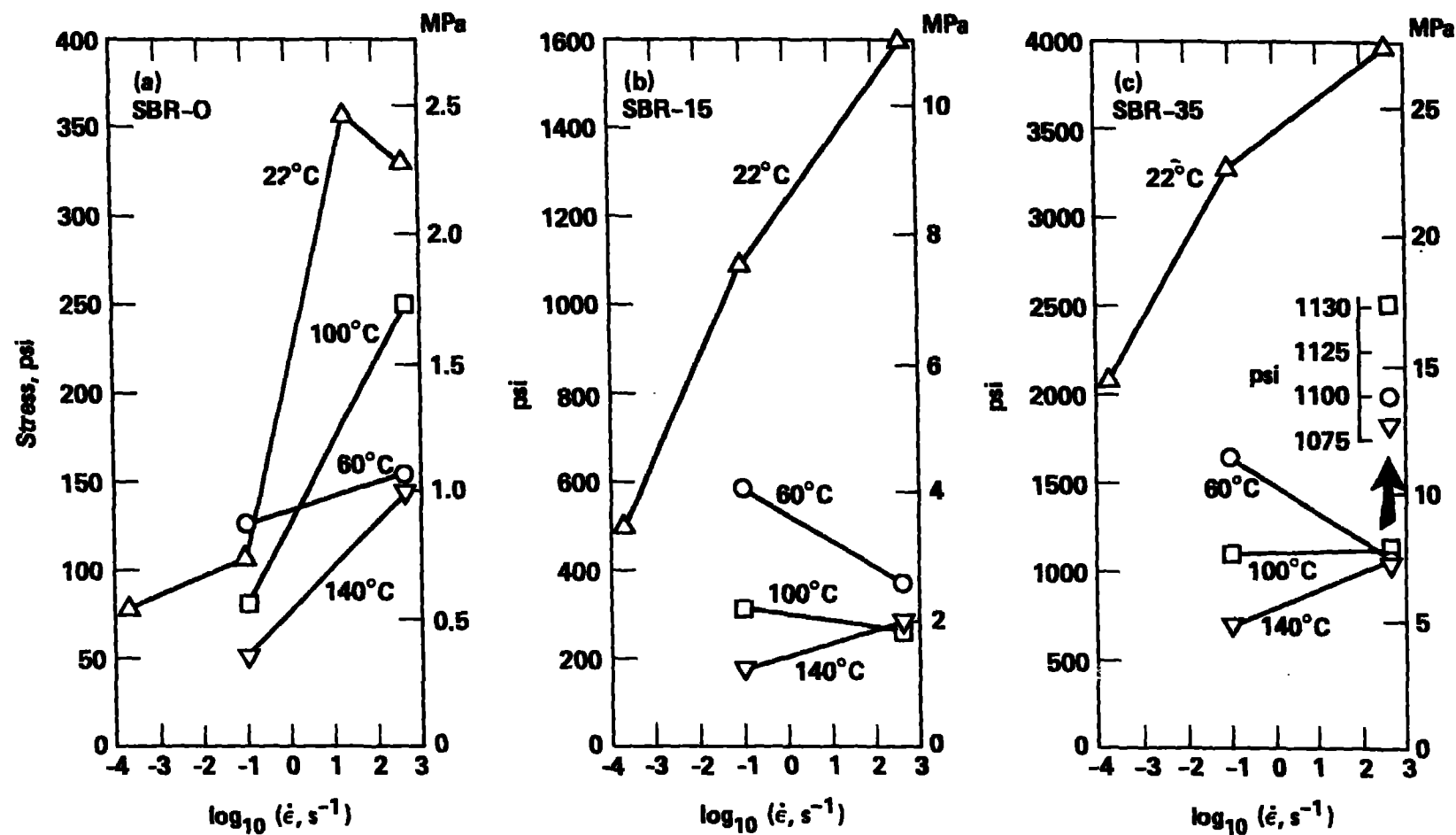


Fig. 24. Effect of strain rate on failure stress or maximum recorded stress at various temperatures for un-nicked strip samples of SBR-0, -15, and -35. NOTE: A number of the SBR-15 and SBR-35 samples could not be taken to failure at the high strain rate.

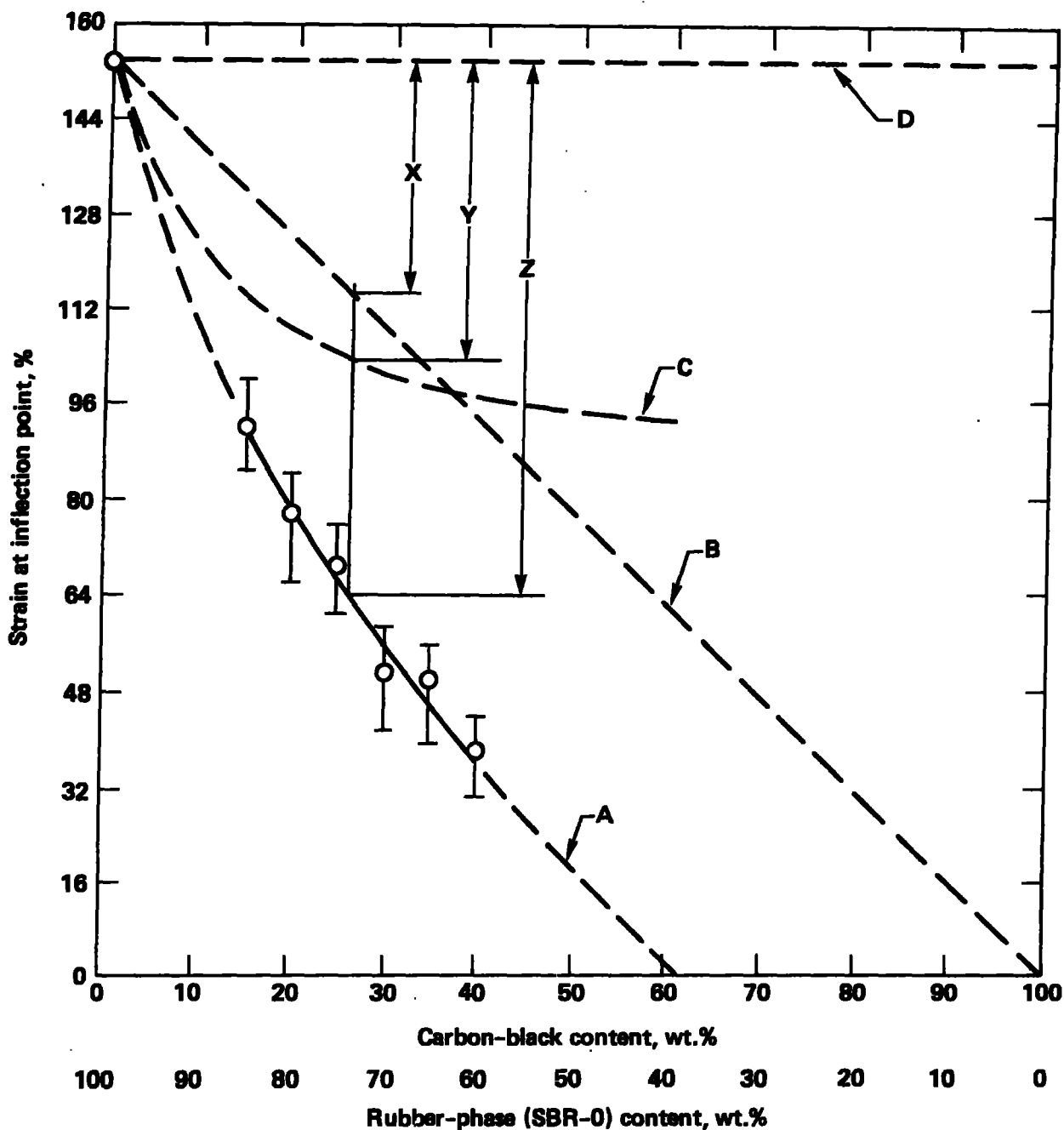


Fig. 25. Effect of filler content on tensile strain at inflection point along the engineering stress-strain curves obtained at a crosshead rate of 0.33 in./s over the temperature range of -20 to 140° C. Curve A refers to experimental data with points representing average values and bars the spread in values; Curve B refers to the effect of rubber phase dilution; Curve C refers to filler-rubber interactions; Curve D refers to a minimum observed inflection strain for SBR-0; $Z = X + Y$.